Selective Inhibition of N-Methyl-d-aspartate Receptors with GluN2B Subunit Protects β Cells against Stress-Induced Apoptotic Cell Death

Anne Gresch, Héctor Noguera Hurtado, Laura Wörmeyer, Vivien De Luca, Rebekka Wiggers, Guiscard Seebohm, Bernhard Wünsch, and Martina Düfer

Pharmaceutical and Medicinal Chemistry, Department of Pharmacology (A.G., H.N.H., L.W., V.D.L., R.W., M.D.), and Pharmaceutical and Medicinal Chemistry (B.W.), PharmaCampus, University of Münster, Münster, Germany; and Institute for Genetics of Heart Diseases (IfGH), Department of Cardiovascular Medicine, University Hospital Münster, Münster, Germany (G.S.)

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ABSTRACT

Participation of N-methyl-d-aspartate (NMDA) receptors (NMDARs) in the failure of pancreatic β cells during development of type 2 diabetes mellitus is discussed. Our study investigates whether β cell mass and function can be preserved by selectively addressing the GluN2B subunit of the NMDAR. NMDAR activation by NMDA and its coagonist glycine moderately influenced electrical activity and Ca2+ handling in islet cells at a threshold glucose concentration (4–5 mM) without affecting glucose-mediated insulin secretion. Exposure of islet cells to NMDA/glycine or a glucolipotoxic milieu increased apoptosis by 5% and 8%, respectively. The GluN2B-specific NMDAR antagonist WMS-1410 (0.1 and 1 μM) partly protected against this. In addition, WMS-1410 completely prevented the decrease in insulin secretion of about 32% provoked by a 24-hour-treatment with NMDA/glycine. WMS-1410 eliminated NMDA-induced changes in the oxidation status of the islet cells and elevated the sensitivity of intracellular calcium to 15 mM glucose. By contrast, WMS-1410 did not prevent the decline in glucose-stimulated insulin secretion occurring after glucolipotoxic culture. This lack of effect was due to a decrease in insulin content to 18% that obviously could not be compensated by the preservation of cell mass or the higher percentage of insulin release in relation to insulin content. In conclusion, the negative effects of permanent NMDAR activation were effectively counteracted by WMS-1410 as well as the apoptotic cell death induced by high glucose and lipid concentrations. Modulation of NMDARs containing the GluN2B subunit is suggested to preserve β cell mass during development of type 2 diabetes mellitus.

SIGNIFICANCE STATEMENT

Addressing NMDA receptors containing the GluN2B subunit in pancreatic islet cells has the potential to protect the β cell mass that progressively declines during the development of type 2 diabetes. Furthermore, this study shows that harmful effects of permanent NMDAR activation can be effectively counteracted by the compound WMS-1410, a selective modulator for NMDARs containing the GluN2B subunit.

Introduction

The glutamate-gated N-methyl-d-aspartate (NMDA) receptor (NMDAR) plays a central role in the mammalian central nervous system. Among others, it participates in neuronal development, synaptic plasticity, learning, and memory formation. Functional NMDARs are heterotetrameric ion channels containing two obligatory GluN1 subunits combined with GluN2 and/or GluN3 subunits. At least eight splice variants are known for the GluN1 subunit that can associate with different isoforms of GluN2 (A–D) or GluN3 (A or B). The GluN2 subunit plays a very important role for channel characteristics, with important differences among the isoforms (e.g., regarding channel decay rate and thus the time for Ca2+ influx) (Paolelli et al., 2013; Zhang and Luo, 2013). Apart from their physiologic function in development and learning, NMDARs are known mediators of neuronal damage when activated in excess or permanently. GluN2B is thought to be a “bad guy” under conditions fostering excitotoxicity (Mony et al., 2009). Consequently, reducing NMDAR activity by targeting NMDARs with GluN2B subunit is the focus of research for treatment of a variety of neuroinflammatory or neurodegenerative central nervous system disorders, including chronic pain, stroke, Alzheimer, or Parkinson disease (Chazot, 2004).

ABBREVIATIONS: DCF, dichlorodifluorescein; FOPP, fraction of plateau phase; KATP channel, ATP-dependent K+ channel; KRH, Krebs-Ringer-HEPES solution; LXR, liver X receptor; MEA, microelectrode array; NMDA, N-methyl-d-aspartate; NMDAR, NMDA receptor; Ro 25-6981, (3R,3S)-1-(4-hydroxyphenyl)-4-methyl-4-phenylmethyl-1-piperidine propanol; TUNEL, terminal deoxynucleotidyl transferase dUTP nick end labeling; Vm, membrane potential; WMS-1410, 3-(4-phenylbutyl)-2,3,4,5-tetrahydro-1H-3-benzazepine-1,7-diol.
One of the first compounds shown to inhibit GluN2B with high affinity was ifenprodil (Williams, 1993). To improve specificity and to reduce side effects, antagonists with tetrahydro-3-benzazepine-1,7-diol scaffold have been designed by exchanging the piperidine ring of the lead compound to a benzazepine (Tewes et al., 2010a). The compound WMS-1410 (3-(4-phenylbutyl)-2,3,4,5-tetrahydro-1H-3-benzazepine-1,7-diol) (Fig. 6A) represents the most promising GluN2B antagonist of this series. WMS-1410 is characterized by a high selectivity, an increased half-life compared with ifenprodil, and good in vivo tolerability (Tewes et al., 2010a). Since neurons and \( \beta \) cells share characteristic features (e.g., the close coupling of exocytosis of secretory granules to electrical activity [Arntfield and van der Kooy, 2011]), it is not surprising that NMDARs are expressed in pancreatic islets. GluN1 and GluN2 subunits were detected on the level of mRNA and protein in insulin-secreting cell lines and rodent islets (Gonoi et al., 1994; Inagaki et al., 1995; Atouf et al., 1997; Morley et al., 2000). The NMDAR is assigned a physiologic role within a negative feedback loop controlling pulsatile insulin secretion (Takahashi et al., 2019) as well as a pathophysiological impact during chronic stimulation by high glucose concentrations (Huang et al., 2017a). Up to now, nothing is known about subtype-specific modulation of the NMDAR and its significance for \( \beta \) cell regulation.

The aim of our study was to elucidate whether inhibition of NMDAR via the GluN2B subunit is suited to interfere with \( \beta \) cell signaling and viability under conditions of increased substrate supply. To address this issue, Ca\(^{2+}\) signaling, electrical activity, insulin release, oxidative status, and apoptotic cell death were determined in mouse islets or islet cells, and the 3-benzazepine derivative WMS-1410 was used for selective targeting of GluN2B.

### Materials and Methods

**Cell and Islet Preparation.** Experiments were performed with islets of Langerhans or single islet cells from adult male and female C57BL/6N mice (Charles River, Sulzfeld, Germany and own breeding, Institute of Pharmaceutical and Medicinal Chemistry, Münster, Germany). The principles of laboratory animal care were followed according to German laws (Az. 53.5.32.7.1/7/2012/668, health and veterinary office Münster, Germany). Mice were euthanized using CO\(_2\), and pancreatic islets were isolated by collagenase digestion. Islets were dispersed to single cells by trypsin treatment. Islets or cells were cultured in RPMI 1640 medium (11 mM glucose) supplemented with 10% fetal calf serum, 100 U/ml penicillin, and 100 \( \mu \)g/ml streptomycin at 37\(^\circ\)C in 5% CO\(_2\) humidified atmosphere. After preparation, islets or dispersed islet cells were kept overnight in standard culture medium. Next day, medium was replaced by either standard medium ± indicated glucose concentrations and substances (only in experiments with acute exposure to the test compounds). Insulin concentration was quantified by a radioimmunnoassay using rat insulin as standard.

**Measurement of Redox Status and \([Ca^{2+}]_o\).** Islets or dispersed islet cells were cultured for 1–2 days on glass coverslips. For islets, glass coverslips were coated with poly-l-lysine. Islets or isolated cells were loaded in bath solution with fura-2 acetoxyethylster (5 \( \mu \)M, 15 mM glucose ± indicated substances, 37\(^\circ\)C, 30 minutes) or 2,7-dichlorodihydrofluorescein (DCHDF)-diacetate (20 \( \mu \)M, 10 mM glucose, 37\(^\circ\)C, 15 minutes) and thereafter maintained in bath solution. For determination of dichlorofluorescein (DCF) fluorescence and \([Ca^{2+}]_o\), after preincubation, the test compounds were omitted from the bath solution during the recordings. Fluorescence was excited at 340 nm and 380 nm every 3 seconds (\([Ca^{2+}]_o\)) or at 480 nm for three consecutive images in 50-millisecond intervals (DCF), and emission was measured by a digital camera (filter 515 nm). Alterations in mean \([Ca^{2+}]_o\) were evaluated by calculating the average of the mean \([Ca^{2+}]_o\) concentration in bath solution with 15 mM glucose for 15 minutes before glucose concentration was changed. To confirm that the islets or islet cells were metabolically intact, bath solution with 0.5 or 15 mM glucose was applied at the beginning or at the end of each experiment. Pancreatic \( \beta \) cells exhibit characteristic glucose-evoked oscillations, which disappear in the presence of a low glucose concentration (Grapengiesser et al., 1988). For determination of DCF fluorescence, two culture dishes were prepared for each condition per mouse preparation. DCF fluorescence was analyzed in two randomly chosen areas of each culture dish by averaging three consecutive measurements per cell. All cells per dish were averaged, and one data point represents the mean of the two samples per mouse preparation.

**Electrophysiology.** Patch-clamp experiments were performed with single islet cells. An EPC-10 patch-clamp amplifier (HEKA, Germany, software “Patchmaster” and “Fitmaster”) was used for data acquisition and analysis. Pipettes were pulled from borosilicate glass capillaries (resistance of 3–5 MΩ). \( V_m \) was measured in the current clamp mode and evaluated by determination of the membrane potential for a period of 1 minute before changing bath solution. Cells were identified as /\( \beta \) cells by glucose-dependent appearance of action potentials. Electrical activity of whole islets was determined by...
extracellular membrane potential recordings with microelectrode arrays (MEA2100-system with 60MEA200/30iR-Ti-gr, Multi Channel Systems, Reutlingen, Germany). Islets were cultured for 3–4 days on microelectrode arrays (MEAs) coated with Matrigel. Medium was changed to bath solution with 3 mM glucose at the beginning of each experiment. Thereafter, glucose was elevated to 8 mM, which was followed by addition of test compounds as indicated. Data were low-pass-filtered at 25 Hz and sampled at 1000 Hz (MC-Rack software). For analysis, the fraction of plateau phase (FOPP; i.e., time with bursting activity related to the entire time interval) was calculated at the end of each experimental maneuver for a period of 20 minutes.

Apoptosis Assay. For determination of the percentage of apoptotic cells, a TUNEL (terminal deoxynucleotidyl transferase dUTP nick end labeling) assay (in situ cell death detection kit, fluorescein, Roche Diagnostics) was used. After the specified incubation time, islets cells were washed with phosphate-buffered saline, fixed with paraformaldehyde 3%, and permeabilized (0.1% Triton-X on ice, 2 minutes). The samples were stained with TUNEL reaction mixture for 1 hour, and this was followed by a nuclear staining with Hoechst-33258 for 10 minutes at room temperature in the absence of light. After nuclei staining, samples were excited at 380 nm, and fluorescence of apoptotic cells was detected at 480 nm. Apoptosis was determined by counting the number of TUNEL-positive cells in relation to all cells in 10 randomly selected fields of each sample (300–400 cells for each mouse islet cell preparation).

Statistics. Data were collected from islets or islet cells of at least three independent mouse preparations for each series of experiments. Values are given as scatter plots and mean ± S.D. Data were normally distributed (GraphPad Prism 3.00). The null hypothesis of each series of experiments was investigated. Statistical significance was assessed by paired test (Fig. 3) or unpaired (Supplemental Fig. 1B) Student’s t test or by ANOVA followed by Student-Newman-Keuls post hoc test for multiple comparisons (Figs. 1, 2, 4, and 5; Supplemental Fig. 1A). GraphPad Prism Windows version 3.00 was used for analysis. Values of $P \leq 0.05$ were considered as statistically significant.

Results

Inhibition of NMDARs Containing the GluN2B Subunit Interferes with the Redox Status of Murine Islet Cells and Protects against NMDA-Induced Cell Damage. Constant activation of NMDAR is linked to cell death and oxidative stress. At first, it was tested whether subtype-specific inhibition of activated NMDAR interferes with the general oxidation status of islet cells. To address this question, the ROS-sensitive dye DCDHF was used, and its oxidation toDCF was monitored. Culturing islet cells in standard medium containing the NMDAR coactivator glycine and supplemented with 5 mM NMDA (Huang et al., 2017b) for 48 hours increased the fluorescence of DCF. WMS-1410 had no influence on DCF fluorescence per se but completely prevented the effect of NMDA (Fig. 1A). As these results suggest that targeting the GluN2B subunit of NMDAR makes the islet cells more resistant to oxidative stress, we elucidated whether inhibition of NMDAR by WMS-1410 can protect against apoptosis. Isolated islet cells were treated with NMDA at a concentration of 500 μM and combined with 10 mM glycine in the presence or absence of WMS-1410 for 18 hours. Prolonged activation of NMDAR resulted in a marked increase in apoptotic islet cells (Fig. 1, B and C, gray vs. white bars). WMS-1410 cleared this effect when applied at concentrations of 1 μM (Fig. 1B) or 0.1 μM (Fig. 1C) in parallel with NMDA. To see whether the NMDA-induced reduction of islet cell mass affects insulin release, insulin secretion was determined after 24-hour culture of whole islets with or without NMDA (500 μM) in standard medium containing glycine. Permanent activation of NMDAR decreased insulin release in response to 15 mM glucose (1 hour) by ~30% (Fig. 1D). Importantly, coincubation with 1 μM WMS-1410 completely prevented this effect. Glutamate, the physiologic stimulus of the ionotropic NMDAR, AMPA (α-amino-3-hydroxy-5-methyl-4-isoxazolepropionic acid), and kainate receptors, was also tested for any impact on islet cell viability. This series of experiments showed that the influence of glutamate was highly variable, ranging from no effect to a clear elevation of apoptotic cell death. Overall, the data showed no dose-response correlation (fraction of apoptotic cells after treatment with 0.5 mM glutamate 7 ± 4%, n = 7 vs. 5 mM glutamate: 9 ± 6%, n = 6) and did not culminate in statistically significant cell death even after 7 days of culture (Fig. 1, E and F). Consequently, we could not detect any beneficial effect of WMS-1410 but only a slight tendency with the higher concentration of 1 μM.

WMS-1410 Prevents the Proapoptotic Influence of Glucolipotoxicity. Progressive loss of functional β cells by elevated metabolic stress is a driving force fostering manifestation of type 2 diabetes mellitus. There are several models to mimic excessive nutrient supply in vitro. Lipotoxic conditions can be induced either by pharmacological stimulation of intracellular lipid synthesis or by providing a high extracellular lipid concentration. We used both methods in combination with a supraphysiologic glucose concentration to create a “glucolipotoxic” milieu. The liver X receptor (LXR) agonist T0901317 (10 μM) was chosen to elevate lipid generation, and palmitate (500 μM) was applied to challenge the islet cells from the outside in medium supplemented with 33 mM glucose for a period of 7 days. Both models resulted in a clear rise in the fraction of apoptotic islet cells (Fig. 2, A and B, gray vs. white bars). Coculture with WMS-1410 (1 μM) showed no effect in the LXR-glucolipotoxic model (Fig. 2A). By contrast, WMS-1410 was clearly protective in the palmitate-glucolipotoxic model. The effect amounted to ~30% with 0.1 and ~60% with 1 μM of the GluN2B antagonist WMS-1410 (Fig. 2B). Ro 25-6981 (1 μM), another inhibitor of GluN2B (Fischer et al., 1997), exerted a similar, beneficial effect (Fig. 2C). In line with this, the nonselective and low-affinity NMDAR blocker memantine (Bresink et al., 1996) also reduced the proapoptotic influence of glucolipotoxicity (Supplemental Fig. 1A). As the attenuation of glucolipotoxicity-induced cell death in single islet cells implicates improvement of insulin release, whole islets were cultured in the presence of high glucose and palmitate. In contrast to the hypothesis, coculture with WMS-1410 (1 μM) only resulted in a small, statistically nonsignificant rise in insulin secretion and thus was not able to preserve the secretory response (Fig. 2D). Determination of the insulin content revealed that glucolipotoxicity drastically reduced the insulin content independent of further treatment with WMS-1410 (Fig. 2E).

Analysis of insulin release as percentage of content showed that WMS-1410 increased the exocytotic process, but, obviously, this was not sufficient to outweigh the severe effect of glucolipotoxicity on insulin biosynthesis (Fig. 2F).

Influence of NMDAR on Membrane Potential and Intracellular Ca²⁺ Concentration. As electrical activity and intracellular Ca²⁺ concentration ([Ca²⁺]i) play a central role for regulation of insulin secretion, the influence of NMDA on these parameters was investigated. Upon activation of
NMDAR, Ca\(^{2+}\) influx through these ion channels should lead to a depolarization of the membrane potential and consequently open voltage-dependent Ca\(^{2+}\) channels (Inagaki et al., 1995). On the other hand, there is evidence that NMDAR activation finally triggers a negative feedback via Ca\(^{2+}\)-regulated K\(^{+}\) channels and KATP channels (Marquard et al., 2015). To evaluate this, the pattern of oscillations of the electrical activity was investigated in islets cultured on microelectrode arrays. Stimulation of islets with 8 mM glucose resulted in regular oscillations. Calculation of the FOPP, (i.e., the time when the membrane was depolarized) resulted in an average value of 12%. Acute application of NMDA/glycine (500/10 \(\mu\)M) neither changed the FOPP nor prolonged the electrically silent interburst intervals (Fig. 3A). For a more detailed analysis, we performed patch-clamp experiments in which the influence of NMDA on Ca\(^{2+}\) action potentials and absolute membrane potential was tested. In the perforated-patch configuration wherein cell metabolism is intact, typical Ca\(^{2+}\) action potentials were visible in the presence of 15 mM glucose. After this functional test for metabolic integrity, glucose concentration was lowered from 15 to 5 mM to decrease the number of action potentials close to the threshold for stimulation of electrical activity. This maneuver created conditions in which alterations in both directions could be detected. Thereafter, NMDA/glycine was added to the bath solution. Three out of nine beta cells reacted with an increase in Ca\(^{2+}\) action potentials (Fig. 3B), whereas six cells responded with a membrane potential hyperpolarization (Fig. 3C). To see how NMDA influenced [Ca\(^{2+}\)]\(_{\text{c}}\) when the glucose concentration was lowered below the threshold for Ca\(^{2+}\) oscillations, similar experiments were performed in cells loaded with the Ca\(^{2+}\)-sensitive fluorescence dye fura-2. Beta Cells responded with typical Ca\(^{2+}\) oscillations that disappeared when glucose was lowered from 15 to 5 mM. When NMDA/glycine (500/10 \(\mu\)M) was applied under these conditions, the maneuver was without any effect in 26 out of 34 beta cells, and [Ca\(^{2+}\)]\(_{\text{c}}\) remained low until glucose was

Fig. 1. Targeting the GluN2B subunit of NMDAR reverses the effects of NMDA on DCF fluorescence, apoptosis, and insulin release. (A) Mouse islet cells cultured in standard medium (containing glycine) supplemented with 5 mM NMDA for 48 hours showed a rise in DCF fluorescence indicative of oxidative stress. WMS-1410 (1 \(\mu\)M) protected against this effect. (B) Addition of NMDA (500 \(\mu\)M) and glycine (10 \(\mu\)M) to the buffer solution (KRH) induced a rise in apoptotic cell death that was partly reversed by coapplication of 1 \(\mu\)M WMS-1410 (18 hours). (C) Similar effects were observed with 0.1 \(\mu\)M of the test compound. (D) Insulin release stimulated by 15 mM glucose (1-hour steady-state incubation) was decreased after 24-hour treatment of islets with NMDA (500 \(\mu\)M) in standard medium. The effect was prevented by 1 \(\mu\)M WMS-1410. (E and F) Addition of glutamate (E: 0.5, F: 5 mM) to the culture medium for 7 days tended to elevate apoptosis independent of the presence or absence of WMS-1410 (1 \(\mu\)M). The number of independent mouse preparations (islet cells: A–C, E and F; islets: D) is indicated below the bars. Circles indicate female, triangles male mice. *P ≤ 0.05, ***P = 0.001 as indicated by the bars and in (D) vs. all other conditions.
elevated to 15 mM again (Fig. 4A, black trace in the exemplary recording). In eight islet cells, a $\text{Ca}^{2+}$ rise or a $\text{Ca}^{2+}$ peak could be detected in response to NMDA (Fig. 4A, gray trace). A succeeding set of experiments revealed that this effect of NMDA was prevented by WMS-1410 (1 $\mu$M) in four out of nine cells reacting to NMDA in the washout phase (Fig. 4B).

As outlined before (Fig. 1, A–D), WMS-1410 was highly effective in preventing the negative effects of prolonged exposure of the islet cells and islets, respectively, to NMDA with respect to cell viability and insulin release. Based on the observation that acute application of NMDA was able to elevate $[\text{Ca}^{2+}]_e$ in a subset of islet cells only, the influence of continuous receptor activation on $[\text{Ca}^{2+}]_e$ was investigated. Whole mouse islets were used for this series of experiments. To evoke typical $\text{Ca}^{2+}$ oscillations, the islets were treated with bath solution containing 15 mM glucose (Fig. 4C). The metabolic integrity of the islets was confirmed by switching the glucose concentration from 15 to 0.5 mM glucose at the end of each experiment. Mean $\text{Ca}^{2+}$ (calculated over a period of 15 minutes) in response to 15 mM glucose did not differ between islets cultured with control medium and islets cultured in standard culture medium supplemented with NMDA (500 $\mu$M) for 24 hours. By contrast, the addition of WMS-1410 (1 $\mu$M) to the latter culture condition for 24 hours changed the pattern of oscillations in islets exposed to 15 mM glucose. Oscillations were faster, and $[\text{Ca}^{2+}]_e$ did not return to a basal level between oscillations or remained in a long plateau phase after rising (Fig. 4C, lower vs. upper exemplary recording). Consequently, mean $\text{Ca}^{2+}$ was higher in WMS-1410–preincubated islets (Fig. 4C, diagram). These data suggest that the protective effect of WMS-1410 on the NMDA-mediated lowering of insulin release (Fig. 1D) was achieved by two factors: preserved $\beta$ cell mass and an increased $\text{Ca}^{2+}$ influx in response to glucose stimulation.
Control experiments were performed (Supplemental Fig. 1B) to exclude that WMS-1410 influences \([\text{Ca}^{2+}]_c\) by off-target effects on the KATP channel, which is the most important ion channel coupling glucose metabolism to \([\text{Ca}^{2+}]_c\).

**NMDAR Activation and GluN2B Inhibition Have No Effect on Insulin Release under Physiologic Conditions.** The data presented above illustrated that NMDA had only small effects on membrane potential and \([\text{Ca}^{2+}]_c\) at threshold glucose concentrations and no effect on membrane potential oscillations or \([\text{Ca}^{2+}]_c\) at stimulatory glucose concentrations. These observations were reflected in the lack of any acute effect of NMDA (500 µM, in combination with 10 µM glycine) on insulin release (1-hour steady-state incubation) at glucose concentrations of 6, 8, 10, and 15 mM (Fig. 5A). The combined application of WMS-1410 (1 µM) and NMDA did not change this result (Fig. 5B).

Finally, the influence of WMS-1410 on islets not exposed to any pathologic environment or NMDA was investigated. Isolated islets were treated with 1 µM WMS-1410 for different periods of time (Fig. 5C). As expected, WMS-1410 did not change secretion when added during the 1-hour steady-state incubation with 15 mM glucose. Secretion was also unaffected when the islets were cultured for 7 days in standard medium supplemented with WMS-1410 (1 µM) and thereafter acutely stimulated by 15 mM glucose for 1 hour.

**Discussion**

**β Cell Signaling during Acute versus Continuous NMDAR Activation.** The influence of NMDA and NMDAR activation on insulin secretion was investigated by several groups, but the results are heterogenous (Gonoi et al., 1994; Inagaki et al., 1995; Molnář et al., 1995; Marquard et al., 2015; Patterson et al., 2016; Huang et al., 2017a; Lockridge et al., 2021). The reasons are manifold, including variations in time, concentration of NMDA or glucose, costimuli, or differences in species. In our experiments, acute exposure of the islets to NMDA neither affected insulin release at the threshold for glucose-induced insulin release nor at stimulatory glucose concentrations (Fig. 5A). This lack of effect fits to the microarray data illustrating that the fraction of plateau phase was not altered by NMDA/glycine (Fig. 3A). NMDA also did
not prolong the interburst intervals, an effect one would expect following the idea that—in addition to Ca^{2+} influx via the L-type Ca^{2+} channels—NMDARs interact with and contribute to the negative feedback loop regulating the oscillatory activity of pancreatic islets (Krippeit-Drews et al., 2000; Marquard et al., 2015). Our data suggest that either the NMDAR is not involved in this mechanism or that its contribution due to the irregular distribution (Patterson et al., 2016; Wu et al., 2017) is too small to be detected as a single regulatory component involved in this mechanism or that its contribution due to the low number (11%) of all glucose-responsive islet cells. Currently it is not clear why the fraction of NMDA-responsive cells is rather small. However, as only less than 50% of cells contain additional Ca^{2+} influx via NMDAR at an already existing high level of [Ca^{2+}]_c. To see whether NMDA/glycine-induced effects at substimulatory glucose levels could be monitored by determination of global [Ca^{2+}]_c, we performed experiments in which glucose was lowered slightly below the threshold for Ca^{2+} influx. Treatment with NMDA/glycine induced a Ca^{2+} peak or a sustained, small increase in 24% of all cells (Fig. 4A). This is congruent to data of Inagaki et al. (1995) with rat islets showing an NMDA-induced rise in [Ca^{2+}]_c, in a low number (11%) of all glucose-responsive islet cells. Currently it is not clear why the fraction of NMDA-responsive cells is rather small. However, as only less than 50% of cells contain the mandatory GluN1 subunit, less than half of the cells may express functional NMDAR as suggested for human β cells and for insulin-secreting INS-832/13 cells (Wu et al., 2017). Treatment with the GluN2B-selective compound WMS-1410 further reduced the number of NMDA-responsive islet cells to 15% (Fig. 4B), indicating that NMDARs with this subunit are functionally active in islet cells. Contrasting to the small alterations induced by acute activation of NMDAR, a prolonged, 24-hour incubation with

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**Fig. 4.** Effects of NMDA and WMS-1410 on [Ca^{2+}]_c of islet cells and islets. (A) Islet cells were stimulated with 15 mM glucose to test for metabolic integrity. Thereafter the glucose concentration was lowered until the oscillations stopped. Acute treatment with 500 µM NMDA (+10 µM glycine) induced a rise or a Ca^{2+} peak in 24% of all cells tested. (B) When WMS-1410 (1 µM) was added prior to the NMDA stimulus, the NMDA-induced rise in [Ca^{2+}]_c, was lowered to 15%. Exemplary recordings are presented on the left of (A) and (B). Data are summarized in the pie charts. (C) Twenty-four-hour culture of islets in standard culture medium in the presence of absence of NMDA (500 µM) did not influence the oscillations in [Ca^{2+}]_c evoked by 15 mM glucose (upper trace, dashed vs. continuous line). By contrast, [Ca^{2+}]_c was proportionally shifted to a plateau when the islets were stimulated by 15 mM glucose after 24-hour treatment with WMS-1410 (1 µM) in standard culture medium. Data are summarized in the diagram. The number of experiments with islet cells (A and B) or islets (C) refers to three independent preparations (A and B: females, C: male/female: 1/2). *P ≤ 0.05 vs. all other conditions.
NMDA reduced glucose-mediated insulin release by about 30% (Fig. 1D). This effect was not correlated to any alteration in \([\text{Ca}^{2+}]_{i}\) (Fig. 4C). Our data indicate that the elevated rate of apoptosis might be the reason for the impairment of islet function. Up to now the mechanisms inducing apoptosis remain to be elucidated. It was suggested that mitochondrial damage by oxidative stress as well as induction of endoplasmic reticulum stress play a decisive role (Huang et al., 2019). In agreement with this, we observed a rise in the fluorescence of the ROS-sensitive dye DCF after long-term treatment with a high concentration of NMDA (Fig. 1A). The fact that the \(\text{Ca}^{2+}\) signaling of islets was not impaired after a 24-hour exposure to 500 \(\mu\text{M}\) NMDA argues against a mechanism involving a dramatic decrease in mitochondrial ATP production. Accordingly, determination of the influence of NMDA on ATP content and the ATP to ADP ratio in the mouse insulinoma cell line MIN6 cells showed a decrease by 5 and 10 mM NMDA, whereas 1 mM NMDA was ineffective (Huang et al., 2017b).

**Protective Effects of WMS-1410 on Murine Pancreatic Islets under Stress Conditions.** Our data demonstrate for the first time that blocking NMDAR via GluN2B interacts with apoptotic \(\beta\) cell death. In agreement with the detrimental role of oxidative stress as a trigger for apoptotic pathways (Drews et al., 2010), protection against NMDA-induced cell stress by WMS-1410 reduced the rise in cell death. As the beneficial effect of WMS-1410 on apoptosis was only partial, we tested whether this was enough to counteract the decline in insulin release in response to treatment with NMDA/glycine. Indeed, acute glucose-mediated insulin secretion reached the level of control when GluN2B-containing NMDARs were inhibited during the 24-hour culture of islets with NMDA/glycine (Fig. 1D). Although several reports address the impact of NMDAR antagonists under physiologic conditions, data with diabetic in vivo or in vitro models are scarce. Treatment of streptozotocin-diabetic mice with the NMDAR antagonist memantine attenuated the diabetic phenotype and lowered the rate of caspase3-positive islet cells (Huang et al., 2017a). In leptin receptor–deficient (db/db) mice, 3 mg/ml dextromethorphan in the drinking water ameliorated the time-dependent development of hyperglycemia compared with treatment with 1 mg/ml of the drug (Marquard et al., 2015). This was accompanied by a rise in insulin content. As we have previously shown that dextromethorphan exerts unspecific effects on L-type \(\text{Ca}^{2+}\) and \(\text{K}_{\text{ATP}}\) channels (Gresch and Düfer, 2020), the contribution of NMDAR inhibition to the observed effects remains unclear. As lipids considerably promote \(\beta\) cell damage during development of type 2 diabetes mellitus (Lytrivi et al., 2020), we used two approaches to address the influence of GluN2B-selective inhibition of NMDAR on apoptosis in an environment mimicking the in vivo situation in an in vitro model. On the one hand, high glucose was combined with palmitate, and on the other hand, the LXR activator 70901317 was used to interfere with intracellular lipid handling by fostering intracellular lipid accumulation (Choe et al., 2007). In the high-glucose/palmitate model, WMS-1410 and Ro 25-6981 markedly reduced the rate of apoptosis.
of apoptosis (Fig. 2, B and C). By contrast, the NMDAR-inhibiting approach was not effective in the high-glucose/T0901317 model (Fig. 2A). At a first glance, the latter result might appear contradictory, as T0901317 increases apoptotic cell death similar to the effect observed with palmitate in our experiments with islet cells and was reported to decrease insulin secretion in MIN6 cells (Meng et al., 2012). Meanwhile, it is known that this compound directly inhibits mitochondrial respiration (Maczewsky et al., 2017) and exerts unspecific effects in islet cells of LXRα/β-double-knockout mice (Maczewsky et al., 2020). Consequently, it is not astonishing that inhibition of NMDAR via GluN2B could not prevent this rather unspecific cell damage rendering this model system unsuitable. The experiments with high glucose/palmitate demonstrate that glucose-stimulated insulin release could not be rescued by WMS-1410 for 24 hours. This indicates activation of a compensatory mechanism and might be the reason for the slight trend to an increased secretory response but, obviously, elevated Ca2+ influx is not enough to protect cell function against the manifold effects of glucolipotoxicity.

In summary, this study shows that acute activation of NMDAR does not severely interact with the stimulus-secretion coupling, whereas excessive stimulation of the receptor leads to oxidative stress, fosters cell death, and decreases insulin release. Targeting the GluN2B subunit of NMDAR protects against oxidative stress and decreases the rate of apoptotic cell death observed in the NMDA model and in an in vitro glucolipotoxicity model (Fig. 6B). Further studies with in vivo approaches are necessary to evaluate whether modulators of the GluN2B isoform are suited to support the maintenance of glycemic control in type 2 diabetes mellitus.

**Authorship Contributions**

**Participated in research design:** Gresch, Düfer.

**Conducted experiments:** Gresch, Noguera Hurtado, Wörmeyer, De Luca, Wiggers.

**Contributed new reagents or analytic tools:** Wünsch.

**Performed data analysis:** Gresch, Noguera Hurtado, Wörmeyer, Wiggers, De Luca.

**Wrote or contributed to the writing of the manuscript:** Gresch, Seebohm, Wünsch, Düfer.

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**References**


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Address correspondence to: Dr. Martina Dufer, Pharmaceutical and Medicinal Chemistry, Department of Pharmacology, PharmCampus, University of Münster, Münster 48, 48149 Münster, Germany. E-mail: martina.dufer@uni-muenster.de
Supplementary Figure

Selective Inhibition of NMDA receptors with GluN2B subunit protects beta cells against stress-induced apoptotic cell death

Anne Gresch§, Héctor Noguera Hurtado§, Laura Wörmeyer§, Vivien De Luca§, Rebekka Wiggers§, Guiscard Seebohm§, Bernhard Wünsch§, Martina Düfer§

§ Pharmaceutical and Medicinal Chemistry, Dept. of Pharmacology, PharmaCampus, University of Münster, Corrensstraße 48, 48149 Münster, Germany

Institute for Genetics of Heart Diseases (IfGH), Department of Cardiovascular Medicine, University Hospital Münster, D-48149 Münster, Germany.

§ Pharmaceutical and Medicinal Chemistry, PharmaCampus, University of Münster, Corrensstraße 48, 48149 Münster, Germany
Influence of memantine on glucolipotoxicity-induced apoptosis and effect of WMS-1410 on K<sub>ATP</sub> current. (A) After a culture period of 7 days, apoptotic islet cell death increased in the G33/palmitate-model. The non-subunit-selective NMDAR blocker memantine (1 μM) partially protected against this. (B) K<sub>ATP</sub> current of single beta cells was measured in the standard whole-cell configuration after 7 days of incubation with control medium ± WMS-1410 (1 μM). The GluN2B-specific inhibitor did not affect the current. Recordings were performed with bath solution (0,5 mM glucose, see methods section of the main document). Pipette solution contained (in mM): 130 KCl, 4 MgCl₂, 2 CaCl₂, 10 EGTA, 0.65 Na₂ATP, 20 HEPES (pH 7.15). K<sub>ATP</sub> current was recorded by application of 300-ms pulses to -60 mV and -80 mV, respectively, from a holding potential of -70 mV. The amplitude at -60 mV was evaluated (mean of 3 consecutive data points) when steady state was reached. At the end of each recording tolbutamide (100 μM) was added to verify the measurement of K<sub>ATP</sub> currents. ***p≤0.001, **p≤0.01, ###p≤0.001 vs. GLT conditions.