Vasopressin increases GAGA binding activity to the V1b receptor promoter through transactivation of the MAP kinase pathway

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Abstract

Previous studies show that binding of nuclear proteins to GAGA repeats (GAGA box) in the vasopressin type 1b receptor (V1bR) promoter is essential for transcriptional initiation of the gene. To determine whether increased vasopressin (VP) during stress activates V1bR expression through the GAGA box, we examined the effects of VP on GAGA binding activity and on the ability of the V1bR promoter to recruit RNA polymerase in the hypothalamic cell line, H32. In chromatin immunoprecipitation assays, VP induced RNA polymerase II recruitment by the wild type V1bR promoter but not by a construct with the major GAGA box deletion. VP (10 min) also increased binding of nuclear proteins to radiolabeled GAGA oligonucleotides in electromobility shift assays. VP-induced GAGA binding activity was potentiated by the protein kinase C inhibitor, calphostin C, and was prevented by the MEK inhibitor, UO126, and the epidermal growth factor receptor (EGFR) inhibitor, AG1478, suggesting that VP activates GAGA binding through transactivation of the EGFR. This was confirmed by western blot experiments showing rapid increases in phospho ERK after incubation with VP, an effect that was potentiated by calphostin C and inhibited by UO12 and AG1478, as well as by the ability of VP to phosphorylate the EGFR. Using receptor selective VP analogs we showed that both V1aR and V1bR subtypes can mediate GAGA binding activation in H32 cells. This study demonstrates that VP stimulates GAGA binding to the V1bR promoter through transactivation of the EGFR and MAP kinase. The data support the hypothesis that VP contributes to pituitary V1bR upregulation during stress through GAGA binding-mediated transcriptional activation.

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Introduction

The nonapeptide, vasopressin (VP), synthesized in parvocellular neurons of the paraventricular nucleus (PVN) of the hypothalamus is an important regulator of pituitary adrenocorticotropic (ACTH) secretion, mainly by potentiating the stimulatory effect of corticotropin releasing hormone (Gillies et al. 1982, Antoni 1993, Aguilera 1994). The expression and secretion of VP into the pituitary portal circulation (Plotsky 1988, Berkenbosch et al. 1989, Engler et al. 1989, de Goeij et al. 1991, Tannahill et al. 1991, Whitnall 1993) as well as the number of VP receptors in the pituitary, increase markedly during the late phase of an acute stress and following chronic stress (Aguilera et al. 1994, Aguilera & Rabadan-Diehl 2000). The effect of VP in the pituitary corticotrope cells are mediated by the vasopressin type 1b receptor (V1bR) subtype, which is coupled to phospholipase C, leading to increases in cytosolic calcium and activation of protein kinase C (PKC) (Jard et al. 1987, Carvallo & Aguilera 1989, Antoni 1993). The parallel changes in V1bR content in the pituitary and corticotrope responsiveness observed in a number of experimental conditions have suggested that V1bR regulation contributes to the adaptation of the hypothalamic–pituitary–adrenal (HPA) axis during stress (Aguilera 1994). Therefore, a better understanding of the mechanisms controlling V1bR expression may prove important for the management of stress-related disorders involving HPA axis disregulation.

It has been shown that the increases in VP binding to pituitary membranes observed 4 h after stress are associated with biphasic changes in V1bR mRNA in which there is an initial decrease by 2 h, followed by a late increase by 4 h. These rapid changes in mRNA expression suggest that rapid activation of V1bR gene transcription is required in order to restore V1bR mRNA levels (Rabadan-Diehl et al. 1995).

We have previously shown that interaction of a pituitary nuclear protein complex with GAGA repeats located near the transcription start point of the V1bR promoter (inverted GAGA box) is critical for transcriptional activation of the gene (Volpi et al. 2002). GAGA repeats were first identified in Drosophila heat shock
protein and histone gene promoters, where they were found to bind a protein called GAGA factor, encoded by the essential Triithorax-like (Trl) gene (Biggin & Tjian 1988b, Gilmour et al. 1989, Soeller et al. 1993, Farkas et al. 1994). In these genes, GAGA factor binding to GAGA repeats has been implicated in chromatin structure and in remodeling and initiation of transcription (Biggin & Tjian 1988, Gilmour et al. 1989, Farkas et al. 1994). Transfection of Drosophila GAGA factor markedly increases V1bR promoter activity, and gel shift assays using pituitary nuclear proteins revealed marked increases in GAGA binding activity during stress. Moreover, the increases in GAGA binding activity were rapid, 30 min after initiation of stress, suggesting that activation of the binding protein is mediated by phosphorylation (Volpi et al. 2002). Since the secretion of VP increases during stress and there is evidence suggesting that VP could be involved in the regulation of its own receptor (Volpi et al. 2004), it is possible that VP induces GAGA binding activation with consequent stimulation of V1bR transcription.

In the present study we tested the hypothesis that VP contributes to the transcriptional regulation of the V1bR by activating the binding of GAGA binding proteins to the V1bR promoter. A problem with these studies has been the lack of availability of a cell line with reasonable endogenous receptor expression. We have previously reported endogenous V1bR expression in the hypothalamic cell line, H32. However, current passages of these cells show very low level of expression of the receptor and we have not been able to obtain consistent results when evaluating changes of endogenous gene expression. For this reason, we sought to examine the effects of VP on RNA polymerase II recruitment by the V1bR promoter as an index of transcriptional activity. In addition, we used gel shift assays to examine the effects of VP on the ability of nuclear proteins to bind to GAGA repeats.

Materials and methods

Cell culture and treatments

The hypothalamic cell line, H32, provided by Dr Joachim Spiess, Goettingen, Germany, was cultured in Dulbecco’s modified Eagle’s medium (DMEM) (Life Technologies, Inc.) containing 10% fetal bovine serum (FBS) (Life Technologies, Inc.), 10% horse serum and 1% penicillin/streptomycin (Life Technologies, Inc.). After 24 h culture in 100 mm plates (15 × 10⁵ per plate), at 37 °C under 5% CO₂/95% air, the medium was changed to serum-free medium containing 0.1% BSA before incubation with VP (10 nM), or human epidural growth factor (hEGF; 200 ng/ml) (R&D Systems Inc., Minneapolis, MN, USA). To determine the signaling pathways and receptor subtype mediating the effects of VP, cells were incubated in the presence and absence of the following: the non-peptide V1bR antagonist, SR144915, 100 nM; the V1a receptor (V1aR) antagonist, SR49059, 100 nM (both provided by Dr C Serradeil-le Gal, Sanofi-Synthlab, Toulouse, France); the epidermal growth factor receptor (EGFR) inhibitor, AG1478, 100 nM (CalBiochem, San Diego, CA, USA); the PKC inhibitor, calphostin C, 1 µM (Biomol Research Lab., Plymouth Meeting, PA, USA); and the MEK inhibitor, UO126, 10 µM (Tocris, Ellisville, MO, USA). The V1bR agonist d[Cha⁴]-arginine vasopressin (d[Cha⁴]AVP), provided by Dr Maurice Manning (Toledo, OH, USA) was used at a concentration of 10 nM. After incubation for the time periods indicated in the Results section and the figure legends, cells were processed for electrophoretic mobility shift assay (EMSA) or western blot analysis of phospho ERK1/2.

Electrophoretic mobility shift assay

Nuclear extracts from H32 cells were prepared using NE-PER nuclear and cytoplasmic extraction reagent (Pierce, Rockford, IL, USA) supplemented with protease and phosphatase inhibitors (Sigma, St Louis, MI, USA) and 2 mM phenylmethylsulfonyl fluoride (Sigma). Double-stranded oligonucleotide probes were synthesized as complementary single strands (Sigma Genosys, The Woodlands, TX, USA), and annealed at 80 °C for 20 min followed by slow cooling to room temperature overnight. Aliquots of 25 pmol GAGA double-stranded oligonucleotides were end-labeled using T4 polynucleotide kinase (New England BioLabs, Inc, Ipswich, MA, USA) and 50 pmol [γ-3²P]ATP (Perkin Elmer Life Sciences, Wellesley, MA, USA) for 45 min at 37 °C, in a total volume of 50 µl. Labeled GAGA probe was purified by centrifugation through a Sephadex G-25 column (Roche Molecular Biochemicals, Indianapolis, IN, USA). The sequences of the probe used for EMSA was GTGACA(GAGA)₃GGGAA (which corresponded to 3GAGA repeats flanked by the 6 nucleotides found upstream and downstream of the GAGA box in the V1bR. The incubation mixture contained 1 µg nuclear extract, 25 mM HEPES pH 7.9, 0.5 mM EDTA, 12.5 mM MgCl₂, 10% glycerol, 1 mM dithiothreitol, 125 mM KCl, 0.05 µg/µl Poly(dIdC) and 1 µg/µl BSA (Sigma) in a total volume of 20 µl. Radiolabeled oligonucleotide (1 ng) was added to the reaction and incubated for 20 min at room temperature. Protein concentrations were determined using the sensitive BCA protein assay (Pierce), before and after adjusting the protein concentration. Reaction products were electrophoresed on a 5% polyacrylamide gel (29:1 acrylamide:bis-acrylamide) in 0.5 X TBE (Trisborate EDTA buffer) for 90 min at room temperature at 150 V. Gels were vacuum-heat dried and bands were visualized.

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H32 cells were transfected with 4 µg V1 bRp830-Luc and V1 bRp830ΔGAGA-Luc using Lipofectamine Plus (Invitrogen Life Technologies, Carlsbad, CA, USA) in 100 mm plates. Twenty-four hours after transfection, cells were changed to 0·1% BSA media for 30 min before incubation with 100 nM VP for 30 min or 1 h. Chromatin immunoprecipitation (ChIP) assays were performed according to the manufacturer’s protocol (Upstate Biotechnology, Lake Placid, NY, USA) with some modifications. Briefly, cells were cross-linked by adding formaldehyde directly to culture medium to a final concentration of 1% and were incubated for 10 min at room temperature. Cells were then washed twice with ice-cold PBS and collected by scraping in 1 ml ice-cold PBS containing protease (Roche Molecular Biochemicals) and phosphatase (Sigma) inhibitor cocktails. Cells were then resuspended in 250 µl SDS lysis buffer containing protease and phosphatase inhibitor cocktails and incubated on ice for 10 min. The lysates were sonicated four times for 10 s (Misonix Incorporated, Farmingdale, NY, USA) to reduce DNA fragment length to approximately 500–2000 base pairs, and subjected to centrifugation for 10 min to remove debris. Supernatants were collected and diluted 10-fold in ChIP dilution buffer containing protease inhibitor cocktail and phosphatase inhibitor cocktail as above. Aliquots (50 µl) of chromatin solution were removed for quantification of the input DNA in the samples before immunoprecipitation. The remaining chromatin solutions were immunocleared with 75 µl salmon sperm DNA/protein A agarose slurry for 1 h at 4 °C with agitation. Immunoprecipitation was performed using 500 µl precleared samples and 10 µl RNA polymerase II specific antibody (Pol II (H224) 200 µg/ml; Santa Cruz Biotech Inc., Santa Cruz, CA, USA), overnight, under agitation at 4 °C. This was followed by the addition of 60 µl salmon sperm DNA/protein A agarose slurry, incubation for 1 h at 4 °C under agitation, and sequential washes with low salt immune complex wash buffer, high salt immune complex wash buffer and LiCl immune complex wash buffer (Upstate Biotechnology) for 3–5 min. After two additional washes in Tris–EDTA (TE) buffer, immunoprecipitated DNA–protein complexes were eluted twice with 1% SDS and 0·1 M NaHCO3. Pooled eluates, as well as the input DNA aliquots from above, were heated at 65 °C for 4 h to overnight in 0·2 M NaCl to reverse the formaldehyde cross-linking. Following incubation at 45 °C for 1 h in 40 µM Tris–HCl (pH 6·5) containing 10 µM EDTA, and 20 µg proteinase K, DNA fragments were purified with phenol:chloroform:isoamylalcohol (25:24:1) and ethanol precipitation, reconstituted in distilled water and subjected to PCR using 2 sets of primers designed to amplify the GAGA box region in the V1bR promoter. The first set of primers, yielding a 430 bp band containing the GAGA box, was: forward primer 5′-CCTTTAGGGCATGCTTCTCAG-3′, reverse primer 5′-TAGGAAACCCAGTGAGGA-3′. PCR for detection of the promoter construct with the GAGA box deletion was performed using the forward primer, 5′-CGGGACTAGCTCATATCTTTC-3′ and the same reverse primer, to yield a 188 bp product. Each cycle consisted of 40 s at 94 °C, 40 s at 58 °C and 40 s at 72 °C, followed by a 10 min extension at 72 °C. The PCR products were separated and visualized in a 2% Tris acetate/EDTA-agarose gel containing ethidium bromide and sized using PCR markers (Promega). The image was captured electronically and the bands were quantified using KODAK 1D Image Analysis software.

Western blot analysis

Cells were cultured in 10-cm plates at 80% confluence, and pre-incubated with vehicle or appropriate signaling inhibitor for 30 min before addition of VP. After 2·5 min incubation with VP, cells were lysed with T-PER tissue protein extraction reagent (Pierce), and 20 µg protein were loaded and separated in a 10% SDS-PAGE. For EGFR, 200 to 250 µg cell protein lysates in 500 µl lysis buffer were immunocleared by incubation for 2 h at 4 °C with 50 µl protein A-agarose beads and centrifuged at 12 000 × g for 20 s. Supernatants were transferred to a new tube and incubated overnight at 4 °C on a rocking platform, after addition of 5 µl EGFR, 1:100 dilution (Cell Signaling Technology, Beverly, MA, USA). Antibody-bound EGFR was immunoprecipitated by addition of 50 µl protein A-agarose beads, incubation for 1 h at 4 °C and centrifugation. Pellets were washed 5 times with lysis buffer before resuspending in SDS sample buffer, denatured for 5 min at 95 °C and loaded into a 4–20% gradient polyacrylamide gel. All steps before resuspension in SDS buffer were performed in the presence of protease and phosphatase inhibitor cocktail. Proteins were transferred from the gel to a PDVF membrane (Amersham Pharmacia Biotech. Piscataway, NJ, USA), incubated with 5% blocking agent in 1 × TBST (TBS plus 0·1% Tween-20) for 1 h and incubated with the anti-phosphorylated ERK or EGFR (Cell Signaling Technology) antibodies at a 1:1000 dilution or phospho-Tyr 1173-EGFR (Santa Cruz, CA, USA) antibody at 1:500 or anti-α tubulin at 1:5000 (Sigma), overnight. After washing in 1 × TBST, membranes were incubated for 1 h with peroxidase-linked anti-rabbit IgG at a 1:25 000 dilution for phospho-ERK, EGFR and phospho-Tyr 1173-EGFR.
or with anti-mouse IgG at a 1:10 000 dilution for 30 min for α tubulin used to correct for protein loading. The intensity of the bands was normalized for α-tubulin to correct for protein loading. Detection of immunoreactive protein bands was performed using ECL Plus reagents (Amersham Pharmacia Biotech) and exposure to BioMax MR film (Kodak, Rochester, NY, USA). Light transmittance was quantified using a computerized imaging system (Imaging Research, St Catherine, Ontario, Canada), using the public domain NIH Image program (developed at the US National Institutes of Health, and available on the internet at: http://rsb.info.nih.gov/nih-image).

Inositol phosphate production

Transiently transfected CHO cells were cultured in 24-well plates and labeled with 2·5 µCi/ml myo-[3H]inositol per well for 24 h, washed with media containing 0·1% BSA and 10 mM LiCl, and then incubated for 15 min under the conditions indicated in the Results section and in the figure legends. Incubations were stopped by addition of one volume of cold stop solution (1 M KOH, 18 mM sodium tetraborate, 3·8 mM EDTA, 7·6 mM NaOH) followed by neutralization with 7·5% HCl. Total inositol phosphates were separated by anion exchange chromatography as previously described (Aguilera et al. 1994) and measured in a liquid scintillation counter.

Measurement of VP receptors

VP receptors were measured by binding of [3H]VP to 30 000 × g membrane fractions prepared from H32 cells pooled from 3 cultures in 10 cm diameter culture plates at 80% confluence. Binding inhibition curves were performed using 50 µg cell membrane protein, 100 000 c.p.m. [3H]VP (Perkin-Elmer Life Science) and increasing concentrations of unlabeled VP or the selective V1bR antagonist SSR149415, or the V1aR antagonist SR49059 (0·1 to 1000 nM), as previously described (Aguilera et al. 1994).

Data analysis

Statistical significance of the differences between groups was calculated by one-way analysis of variance (ANOVA), followed by the Student–Newman–Keuls method for pairwise multiple comparisons. Statistical significance was set at P<0.05. Data are presented as means ± standard error of the mean (S.E.M.) from the values in the number of observations indicated in the Results section or figure legends.

Results

Effect of VP on Pol II recruitment by the V1bR promoter

The ability of the V1bR promoter (wild type or with deletion of the main GAGA box) to recruit RNA polymerase II (Pol II) following incubation of H32 cells with VP. Cells were incubated with VP for 30 or 60 min, 24 h after transfection with V1bR promoter luciferase constructs with (V1bRp-Luc 830) or with deletion of the GAGA box (V1bRp-Luc 830-GAGA). Cross-linked DNA was immunoprecipitated with Pol II antibody and subjected to PCR using primers against the V1bR promoter. The semiquantitative analysis of the PCR bands after subtraction of the blank values and correction for DNA input in three experiments is shown in the lower panel.

Effect of VP on GAGA binding activity

Since previous studies showed transient increases in GAGA binding activity in pituitary nuclear proteins from stressed rats, we sought to examine the ability of...
VP to mediate the effect of stress, using EMSA with nuclear extracts from H32 cells following incubation with VP. The time course of the changes in GAGA binding activity is shown in Fig. 2. Binding to radiolabeled GAGA oligonucleotides increased by 94.7% at 10 min \( (P<0.05, n=3) \) and declined to levels not significantly different from basal by 15 min.

To determine the receptor subtypes present in H32 cells, we examined the effect of subtype specific VP antagonists and a specific V1bR agonist on the ability of VP to induce inositol phosphate formation and to inhibit the binding of \(^{[3]H}\)VP to 30 000 \( \times g \) membrane-rich fractions from H32 cells. In three experiments, incubation of the cells with VP increased total inositol phosphate formation by 2.8-fold \( (P<0.01) \) but only a minor, though significant increase was observed with the V1bR agonist d[Cha\(^4\)]AVP \( (P<0.05) \). While stimulation by d[Cha\(^4\)]AVP was completely prevented by the V1bR antagonist, SSR149415, and was unaffected by the V1aR antagonist, SR49059, the effect of VP was markedly inhibited \( (82.4\%, P<0.01) \) by the V1aR antagonist and only slightly inhibited by the V1bR antagonist \( (25.3\%, P<0.05) \), suggesting that the predominant receptor subtype is the V1bR (Fig. 3A). Supporting this data, in binding studies the V1aR antagonist, SR49059, inhibited 86.5% of the total binding of \(^{[3]H}\)VP to 30 000 \( \times g \) membrane fractions of H32 cells, while the V1bR antagonist inhibited only 31% of the binding (Fig. 3B). The binding inhibition constants for the V1a and V1b receptor antagonists were similar to that of VP, with \( K_i \) values of 0.94 ± 0.3, 1.1 ± 0.2 and 0.89 ± 0.2 nM respectively.

Since the above data show that H32 cells express both V1a and V1b receptors, we used receptor subtype specific analogs to determine the type of receptor mediating the activating effect of VP on GAGA binding. As shown in Fig. 4, the V1bR agonist d[Cha\(^4\)]AVP or VP increased GAGA binding activity to similar levels \( (177.4 \pm 19.4\% \text{ and } 191.5 \pm 23.4\% \text{ respectively}) \). Co-incubation of the cells with the V1bR antagonist, SSR149415, completely prevented the stimulatory effect of the V1bR agonist d[Cha\(^4\)]AVP but reduced the effect of VP by only 26%. The effect of VP was reduced by 33% by the V1aR antagonist.
VP activates GAGA binding through transactivation of the EGFR

To identify the signaling pathways involved in the activation of GAGA binding by VP, cells were incubated with VP in the presence of several signal transduction inhibitors before testing the activity of nuclear extracts by EMSA. Inhibition of PKC (the main signaling pathway of V1 receptors) using the generic PKC inhibitor, calphostin C, did not prevent but potentiated the stimulatory effect of VP on GAGA binding activity. As shown in Fig. 5, calphostin C alone had no effect but enhanced the stimulatory effect of VP from 148.7% to 191.5% of the basal value (P<0.05, n=3). On the other hand, activation of GAGA binding by VP was completely abolished in the presence of the MAP kinase (MEK 1 and 2) inhibitor, UO126, or the EGFR inhibitor, AG1478, with values not significantly different from basal levels (P>0.05 compared to VP-stimulated activity, n=3) (Fig. 5). Direct stimulation of the EGFR by incubation of H32 cells with EGF (200 ng/ml) caused sustained GAGA binding activation with significant increases at 2.5, 10 or 15 min (P<0.02) compared with basal values at all time points (n=3) (Fig. 6). These results suggest that VP can stimulate GAGA binding activity and transcriptional activation of the V1bR by MEK-activation of MAP kinase through transactivation of the EGFR. The ability of VP to transactivate the MAP kinase pathway was demonstrated by Western blot analysis for phospho-ERK1/2 in protein extracts of cells incubated with VP. As shown in Fig. 7A, phospho-ERK levels were undetectable in H32 cells in basal conditions but became apparent following incubation with VP for 10 min. Similar to GAGA binding activity, VP-induced ERK1/2 phosphorylation was significantly enhanced by the generic PKC inhibitor, calphostin C (P<0.05 vs VP, n=3) and was markedly inhibited by the MEK1/2 inhibitor UO126, (P<0.01, n=3) and the EGFR inhibitor, AG1478 (P<0.01, n=3) (Fig. 7A). To confirm that VP induces ERK1/2 phosphorylation through the EGFR, we examined the effects of VP on EGFR phosphorylation. Western blot analysis of H32 cell proteins using the EGFR antibody after immunoprecipitation with the EGFR antibody revealed a major band of approximately 178 kDa, with molecular size consistent with that of the EGFR (Fig. 7B). A weak band at the same position was obtained using an antibody against the phospho-Tyr1173 of the EGFR. As shown in Fig. 7B and C, incubation of the cells with VP for 5 min had no significant effect on total EGFR content but increased markedly (4.3-fold) the phosphorylated receptor (P<0.01, n=3). Preincubation of the cells for 5 min with the V1 receptor peptide antagonist, Des-Gly[Phaa]1, d[Cha4]AVP (VP or V1bR agonist d[Cha4]AVP for 10 min, with or without 30 min pretreatment with the V1a or V1b receptor antagonists, SR49059 and SSR149415 respectively, to GAGA oligonucleotides is depicted above the bars. Quantification of the intensity of the shifted bands was performed using a Molecular Dynamics PhorImager. The graph shows the mean and s.e. of the values in 3 experiments. *P<0.05 compared with basal (vehicle treated); # P<0.05 compared with VP-stimulated value.

Figure 4 V1 receptor subtype specificity of the effect of VP on GAGA binding activity in H32 cells. EMSA showing the binding of nuclear proteins from H32 cells incubated with VP (10 min) or the V1bR agonist d[Cha4]AVP for 10 min, with or without 30 min pretreatment with the V1a or V1b receptor antagonists, SR49059 and SSR149415 respectively, to GAGA oligonucleotides is depicted above the bars. Quantification of the intensity of the shifted bands was performed using a Molecular Dynamics PhorImager. The graph shows the mean and s.e. of the values in 3 experiments. *P<0.05 compared with basal (vehicle treated); # P<0.05 compared with VP-stimulated value.

Figure 5 Effect of signaling transduction inhibitors on VP-stimulated GAGA binding activity. EMSA using nuclear proteins from H32 cells incubated for 30 min with or without UO126 (MEK1/2 inhibitor), calphostin C (Cal C; PKC inhibitor) or AG1478 (EGFR inhibitor) prior to 10 min exposure to VP, and GAGA oligonucleotides is depicted above the bars. Quantification of the intensity of the shifted bands was performed in a Molecular Dynamics PhorImager. The graph shows the mean and s.e. of the values obtained in three experiments. *P<0.05 higher than basal (vehicle treated); # P<0.05 lower than VP.
d-Tyr(Et)²,Lys⁶,Arg⁸]VP, completely prevented the effect of VP on EGFR phosphorylation.

**Discussion**

The present study demonstrates that VP increases the binding activity of nuclear proteins to GAGA repeats in the V1bR promoter through activation of the MEK MAP kinase pathway. The data also provide evidence that activation of GAGA binding to the V1bR promoter by VP could mediate transcriptional activation of the V1bR gene. Previous studies showing an association between high VP expression in parvocellular neurons of the PVN and pituitary V1bR upregulation during stress, have suggested that VP can upregulate its own receptor (Volpi et al. 2004). This increase in pituitary V1bRs during stress involves increases in V1bR mRNA and, at least in part, activation of the V1bR gene transcription. For example, acute stress causes biphasic changes in V1bR mRNA levels, with an initial decrease by 2 h followed by recovery or elevation above the basal values by 4 h (Rabadan-Diehl et al. 1995). This rapid restoration of V1bR mRNA levels must involve transcriptional activation of the V1bR gene. The present experiments demonstrate that VP can induce recruitment of RNA polymerase II by the V1bR promoter. Since RNA polymerase recruitment is closely associated with initiation of transcription (Kornberg 1999), these data provide strong support for the hypothesis that VP induces V1bR expression through transcriptional activation. Since stress causes rapid release of VP from the median eminence into the pituitary portal circulation (de Goeij et al. 1991), it is likely that the peptide partially mediates the pituitary V1bR upregulation observed during prolonged stress.

Using reporter genes, we previously demonstrated that binding of a nuclear protein complex to GAGA repeats (GAGA box) in the rat V1bR promoter is essential for transcriptional activation of the V1bR gene (Volpi et al. 2002). The present study supports this concept and provides evidence that activation of V1bR transcription by VP is mediated by the GAGA box. This was first demonstrated by the marked reduction in VP-induced RNA polymerase II recruitment by the V1bR promoter construct lacking the GAGA repeats. Secondly, the ability of VP to enhance GAGA binding activity of nuclear extracts of H32 cells suggests that VP stimulates V1bR transcription through activation of
protein binding to the GAGA repeats of the V1bR promoter. While the neuronal cell line, H32, expresses both V1aR and V1bR mRNA, the poor inhibitory effect of the selective V1bR antagonist (compared with the V1aR antagonist) on [3H]VP binding and VP-stimulated inositol phosphate formation indicate that the V1aR is the predominant functional receptor in these cells. The present experiments using receptor subtype specific analogs clearly demonstrate that both V1a and V1b receptors can mediate the stimulatory effect of VP on GAGA binding activity. In spite of the low levels of V1b receptors, the V1bR agonist, d[Cha4]AVP, used at a concentration known to have no interaction with V1aR (Derick et al. 2002), had an effect similar to that of VP, presumably by interacting with the V1bR. This suggests that occupancy of a small number of receptors is sufficient to elicit full activation of GAGA binding. It is also possible that physical interaction between V1bR and V1αR amplifies the V1bR signal. In this regard it has been shown that a number of G protein-coupled receptors, including V1 receptors, can form homodimers and heterodimers (Terrillon 2003, Robert 2005, Young & Aguilera 2005).

The ability of pituitary nuclear proteins to bind GAGA repeats increases during stress and precedes the increases in V1bR mRNA (Volpi et al. 2002). This rapid and transient increase in pituitary GAGA binding activity is consistent with the hypothesis that the binding of a nuclear protein complex to DNA is activated by phosphorylation. Since this protein complex has been only partially characterized (Volpi et al. 2002), we used gel shift assays with GAGA oligonucleotides and signaling transduction inhibitors to identify signaling pathways mediating activation of GAGA binding proteins by VP. Since PKC is the major mediator of the action of VP in pituitary corticotropes and brain (Bilezikjian et al. 1987, Carvallo & Aguilera 1989), the fact that the PKC inhibitor, calphostin C, did not inhibit but actually enhanced VP-stimulated GAGA binding activity in the neuronal cell line, H32, was unexpected. The stimulatory effect of the PKC inhibitor suggests that PKC inhibits GAGA binding activity and that an alternative pathway is responsible for the activating effect of VP on GAGA binding protein. A number of G protein-coupled receptors, including V1 receptors, have been shown to initiate EGF or other growth factor receptor signaling in addition to the G protein-dependent pathways (Leserer et al. 2000, Ghosh et al. 2001, Chiu 2002, Ferguson 2003, Luttrell 2003, Werry et al. 2005). Such a transactivation of growth factor receptors leads to activation of the MAP kinase pathway and in many instances phosphorylation of nuclear proteins (Luttrell 2003). The present demonstration that the MEK inhibitor, UO126, prevented VP-induced activation of GAGA binding indicates that VP stimulates the binding of nuclear proteins to the V1bR promoter through MAP kinase-dependent phosphorylation. In addition, the experiments confirm that VP does indeed induce ERK1/2 phosphorylation in the H32 cell line. The fact that the effects of VP, both on ERK phosphorylation and GAGA binding activation, were blocked by the EGFR inhibitor and were mimicked by EGF suggests that MAP kinase activation by VP requires the involvement of the EGFR. The present demonstration that VP causes rapid phosphorylation of the EGFR, an effect that is prevented by a non-selective V1 VP receptor, confirms the fact that VP causes transactivation of the EGFR and supports an essential role of the EGFR on activation of the MAP kinase pathway by VP.

A GAGA binding protein has been implicated in the transcriptional regulation of vertebrate genes, such as the human type 1 angiotensin II receptor (AT1R), the rat serine protease inhibitor 1, and the Xenopus

**Figure 8** Diagram of the proposed mechanism of transcriptional activation of the V1bR by VP. Activation of V1bR signaling by VP released into the pituitary portal circulation during stress involves transactivation of the EGFR and MAP kinase pathway, leading to phosphorylation and activation of the binding of a nuclear protein capable of interacting with GAGA repeats in the V1bR promoter and activating transcription VPR, vasopressin receptor; PLC, phospholipase C; Gαq/11, alpha subunit of guanyl nucleotide binding protein q/11; PI, phosphoinositides; DAG, diacyl glycerol; PKC, protein kinase C; IP3, inositol triphosphate; EGFR, epidermal growth factor receptor; RAS, monomeric guanyl nucleotide binding protein interacting with RAF; RAF, MAP kinase kinase kinase Raf; MEK, MAP/ERK (extracellular signal regulated kinase) kinase; MAPK, mitogen activated protein kinase.
stromelysin-3 gene (Li et al. 1998, Simar-Blanchet et al. 1998, Wyse et al. 2000). For the AT1 receptor and serine protease inhibitor genes, it has been shown that GAGA binding to the promoter is required for transcriptional initiation (Leverrier 2000). This is consistent with previous studies using reporter genes showing that the GAGA box is required for basal V1bR promoter activity (Volpi et al. 2002), and the present data showing rapid increases in GAGA binding activity, following exposure of the cells to VP, preceding Pol II recruitment by the V1bR promoter. In the case of the serine protease inhibitor 1 and the AT1R, GAGA binding activity is stimulated by growth factors, and the GAGA box appears to mediate growth factor-mediated gene transcription (Simar-Blanchet et al. 1998, Wyse et al. 2000). VP has recognized effects as a growth factor and activation of the V1aR can transactivate the EGFR in a number of tissues, including kidney mesangial cells and intestinal epithelial cells (Ghosh et al. 2001, Chiu 2002).

In summary, the present demonstration that VP induces recruitment of RNA polymerase II by the V1bR promoter and the levels of V1bR mRNA, supports the view that VP exerts a positive regulatory effect on V1bR expression. As shown in Fig. 8, this effect of VP involves activation of the binding of nuclear proteins to GAGA repeats in the V1bR promoter. This effect is not mediated by protein kinase C but by transactivation of the EGFR and MAP kinase. The data provide a mechanism by which increased parvocellular VP expression can activate V1bR transcription and contribute to the pituitary V1bR upregulation observed during stress.

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