



Human protein arginine methyltransferases (PRMTs) can be optimally active under nonphysiological conditions

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Protein arginine methylation is involved in many biological processes and can be enhanced in cancer. In mammals, these reactions are catalyzed on multiple substrates by a family of nine protein arginine methyltransferases (PRMTs). However, conditions that may regulate the activity of each enzyme and that may help us understand the physiological role of PRMTs have not been fully established. Previous studies had suggested unexpected effects of temperature and ionic strength on PRMT7 activity. Here we examine in detail the effects of temperature, pH, and ionic strength on recombinant human PRMT1, PRMT5, and PRMT7. We confirmed the unusual temperature dependence of PRMT7, where optimal activity was observed at 15 °C. On the other hand, we found that PRMT1 and PRMT5 are most active near physiological temperatures of 37 °C. However, we showed all three enzymes still have significant activity at 0 °C. Furthermore, we determined that PRMT1 is most active at a pH of about 7.7, while PRMT5 activity is not dependent on pH in the range of 6.5 to 8.5. Significantly, PRMT7 is most active at an alkaline pH of 8.5 but shows little activity at the physiological intracellular pH of about 7.2. We also detected decreased activity at physiological salt conditions for PRMT1, PRMT5, and PRMT7. We demonstrate that the loss of activity is due to the increasing ionic strength. Taken together, these results open the possibility that PRMTs respond in cells undergoing temperature, salt, or pH stress and demonstrate the potential for *in vivo* regulation of protein arginine methylation.

Protein diversity heavily relies on many enzymes that modify amino acid residues. A major site of protein posttranslational modification is methylation at the guanidino group of arginine residues. These modifications create distinct interactions between proteins and play a significant role in biology including transcription of DNA, activation of enzyme activity, and protein degradation (1–4). In mammals, nine specific enzymes (protein arginine methyltransferases [PRMTs]) have been characterized and are classified based on the specific reaction they catalyze (5–7). While all PRMTs can initially catalyze monomethylarginine formation, type I PRMTs (PRMT1, 2, 3, 4, 6, and 8) catalyze asymmetric dimethylarginine formation and type II PRMTs

(PRMT 5 and 9) catalyze symmetric dimethylarginine formation, while type III PRMTs (PRMT7) only catalyzes monomethylarginine formation. Determination of the *in vitro* activity of these purified PRMT enzymes can give important clues to their substrate specificity and their possible regulation (8–12).

For intracellular cytosolic mammalian enzymes, the assumption is usually made that activity will be at or near optimum levels at the physiological pH of about 7.2 (13), an ionic strength of about 120 mM (14), and a temperature of 37 °C (15). However, we were surprised to find that PRMT7 had little activity at 37 °C and maximum activity was noted at colder temperatures (16, 17). Additionally, little activity was found at the physiological salt concentrations that occur within a cell (16). However, these measurements were generally not taken under initial velocity conditions and we were thus interested in a more detailed picture of PRMT7 catalysis, especially under nonphysiological conditions. We were also interested in comparing the catalysis shown by PRMT7 with that of PRMT1, the major type I enzyme, creating asymmetric dimethylarginine, and PRMT5, the major type II enzyme, creating symmetric dimethylarginine residues.

Here we show that human PRMT1, PRMT5, and PRMT7 activity is optimal under varying conditions. Specifically, PRMT7 activity is optimal at subphysiological temperatures in an alkaline environment containing low salt concentrations. We have also shown that PRMT1 and PRMT5 activities are most optimal closer to physiological temperature and pH, but where reaction mixtures contain little salt.

Results

GST-HsPRMT7 is active at physiological temperatures but has optimal activity at subphysiological temperatures

Recent findings have demonstrated that human histone H2B (HsH2B) is an excellent substrate for PRMT7, with methylation sites detected at arginine residues 29, 31, and 33 (10). A synthetic peptide containing residues 23 to 37 (HsH2B (23–37)) was also an excellent substrate (10). Significantly, optimal PRMT7 activity was found at about 20 °C with very little activity at 37 °C (16). Similar effects were found with the bacterially expressed GST human enzyme fusion protein and the insect-expressed untagged mouse enzyme (10). However, because these activities were measured at long incubation times, we wanted to determine the temperature dependence

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Stress and protein arginine methyltransferase enzymology

under initial rate conditions. We first determined that GST-HsPRMT7 activity is linear with time up to 1 h at both 20 °C and 37 °C with the HsH2B (23–37) peptide (Fig. 1). At 37 °C, we found only about 25% of the activity as found at 20 °C. We also showed no significant increase of activity when the peptide concentration was raised from 10 μ M to 50 μ M, indicating that enzyme was saturated with the peptide substrate at the lower concentration. These results now clearly show that PRMT7 activity is not optimal at the physiological temperature.

Similar results to those shown in Figure 1 with the HsH2B (23–37) peptide substrate were found when we used the GST-GAR protein substrate (Fig. 2). At the 60-min and 120-min timepoints, only about 20 to 30% of the activity at 20 °C was found at 37 °C. Using an alternate assay where the 3 H-methylated GST-GAR polypeptides formed in a 2-h assay were fractionated by SDS-PAGE and analyzed by densitometry following fluorimetry, we found 9.2% of the activity at 37 °C as found at 20 °C (data not shown). We also used this latter assay with a recombinant full-length human HsH2B protein and found less than 11% of the activity at 37 °C as found at 20 °C (data not shown).

In Figure 3, we measured GST-HsPRMT7 activity at temperature values between 0 °C and 45 °C with the HsH2B (23–37) peptide using the P81 phosphocellulose assay. We found the peak of activity at 15 °C with half activity at 0 °C and at 20 to 25 °C. At the physiological temperature of 37 °C, only about 20% of the activity was found compared to that of 15 °C. These data clearly show that PRMT7 is most active in the cold.

Decreased human PRMT7 activity at 37 °C is not due to enzyme instability

To determine if incubation at 37 °C irreversibly denatured PRMT7, we performed the experiment in Figure 4. Here, we preincubated the enzyme at 4 °C, 20 °C, and 37 °C for 2 h and

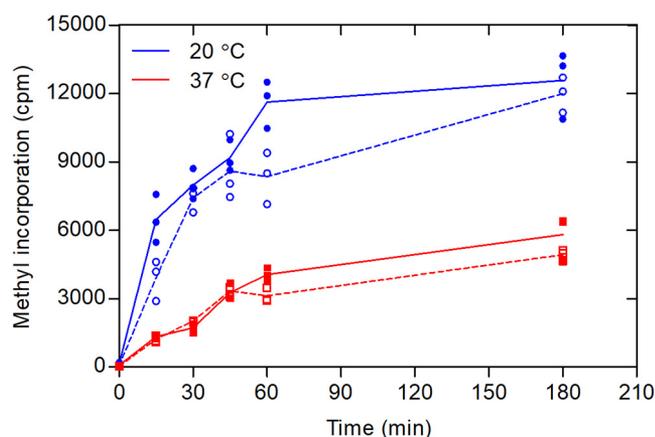


Figure 1. Human PRMT7 is more active at subphysiological temperatures with histone H2B peptide as a substrate. P81 phosphocellulose assay of GST-HsPRMT7 and HsH2B (23–37) was performed at 20 °C and 37 °C as described in the [Experimental procedures](#) section. Five micrograms of GST-HsPRMT7 was incubated with peptide and 0.14 μ M [3 H]AdoMet. *Blue lines* represent reaction mixtures at 20 °C using 10 μ M peptide (*closed circles*) or 50 μ M peptide (*open circles*). The *red line* represents reaction mixtures at 37 °C using 10 μ M peptide (*closed squares*) or 50 μ M peptide (*open squares*). HsH2B, histone H2B; PRMT, protein arginine methyltransferase.

then assayed activity at each of these temperatures. We found little or no loss of activity when assayed at 4 °C, suggesting that PRMT7 was not denatured at 37 °C. In all cases, similar activities were found when each of the incubated samples was assayed at the indicated temperature. Additionally, previous work had shown that the circular dichroism spectrum of PRMT7 was identical at 4 °C, 17 °C, and 37 °C, suggesting that the enzyme is stable at these temperature ranges (18). Additionally, it was shown that the molecular weight of PRMT7 by sedimentation equilibrium analysis was also identical at these three temperatures (18).

His-HsPRMT1 and HsPRMT5/MEP50 are most active near physiological temperatures

We then wanted to establish the generality of the methyltransferase activity at subphysiological temperatures with the major type I enzyme PRMT1 and the major type II enzyme PRMT5. We thus tested His-HsPRMT1 and HsPRMT5/MEP50 under similar conditions to those used above for GST-HsPRMT7 (Fig. 5). For PRMT1, we found a broad optimal peak between 20 °C and 37 °C with half optimal activities at about 15 °C and 45 °C. For PRMT5, the temperature optimum was found at 37 °C with half optimal activity of 25 °C and over 45 °C. Taken together, these results show that PRMT1 and PRMT5 have temperature optimums near physiological body temperature and show distinct patterns from PRMT7. However, we did detect the activity of both enzymes throughout this temperature range.

PRMT1, PRMT5, and PRMT7 are all active at 0 °C

We found that PRMT1, PRMT5, and PRMT7 all displayed activity at 0 °C. To confirm this surprising result, we performed additional time course experiments in assays done entirely on ice (Fig. 6). All of these enzymes were found to have significant activity at 0 °C. GST-HsPRMT7 activity was almost 6-fold higher at 0 °C than at 37 °C. His-HsPRMT1 activity was 6-fold lower in activity at 0 °C than in reaction mixtures that were performed at 37 °C, and HsPRMT5/MEP50 activity was about 3-fold lower at 0 °C than at 37 °C.

PRMT7 is active at alkaline pH

Our finding that PRMT7 is most active at nonphysiological temperatures lead us to examine its activity as a function of pH. As shown in Figure 7A, we were surprised to find little to no activity at the physiological pH of 7.2. This enzyme was found to be more active at alkaline pH values with a pH optimum of 8.4. We performed a control experiment showing that changes in pH did not affect the P81 phosphocellulose assay (Fig. S1). These results suggest that PRMT7 might respond to changes in cellular pH that lead to alkalization.

We also performed pH studies using a substrate of GST-GAR and an SDS-PAGE assay as described in the [Experimental procedures](#) (Fig. S2). The results shown here are very similar to those found in Figure 7A with the HsH2B (23–37) peptide and the P81 phosphocellulose assay.

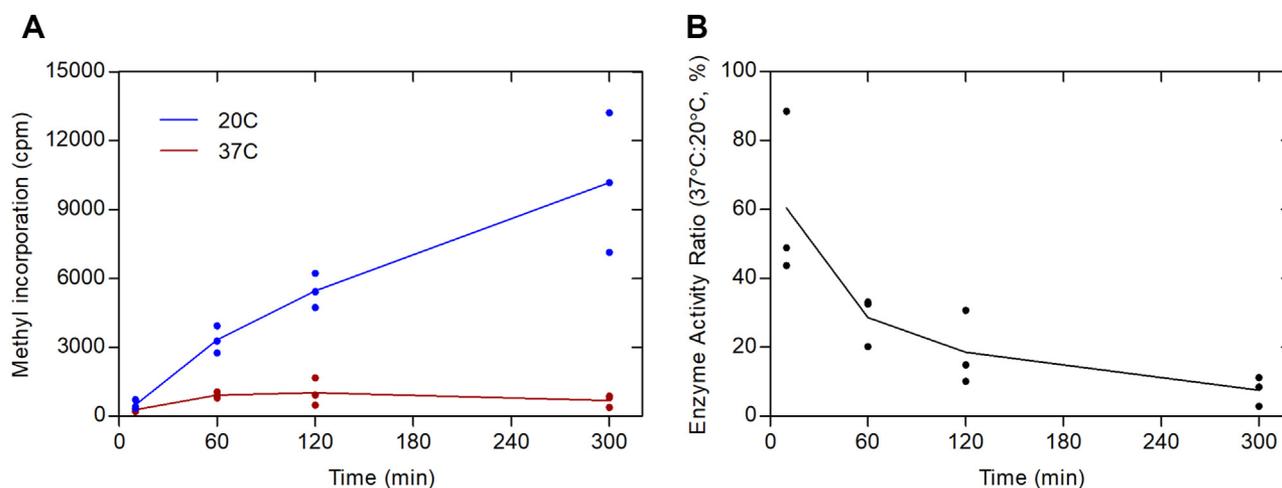


Figure 2. Human PRMT7 is more active at subphysiological temperatures with GST-GAR as a substrate. A, P81 phosphocellulose assay of GST-HsPRMT7 activity. GST-HsPRMT7, 6.2 μ g GST-GAR, and 0.7 μ M [3 H]AdoMet was incubated at either 20 $^{\circ}$ C or 37 $^{\circ}$ C as described in [Experimental procedures](#). The blue line represents the averages of individual reactions at 20 $^{\circ}$ C (blue dots). The red line represents individual replicates of reaction mixtures tested at 37 $^{\circ}$ C (red dots). B, ratios of reactions performed at 37 $^{\circ}$ C to 20 $^{\circ}$ C. The black line represents the averages of the individual ratios calculated from A. PRMT, protein arginine methyltransferase.

PRMT1 and PRMT5 have distinct pH activity profiles from PRMT7

To compare the activity of the major PRMT1 type I enzyme and the major PRMT5 type II enzyme with PRMT7 as a function of pH, we performed the assays shown in [Figure 7, B](#) and C. We found that His-PRMT1 has a pH optimum of about 7.7 with good activity at the physiological pH of 7.2. Additionally, we found that PRMT5/MEP50 is equally active across a wide range of pH values between 6.6 and 8.5. Here, PRMT1 and PRMT5 appeared to differ from PRMT7 in showing activity at the physiological pH.

PRMT7 is inhibited by the addition of salts

Previous studies have shown that GST-HsPRMT7 activity decreases with increasing NaCl concentrations with little

activity seen at physiological salt concentrations, particularly for the HsH2B (23–37) peptide (16). Additionally, a loss of approximately 50% activity was found in 2 mM magnesium and calcium salts (16). Given our results above that PRMT7 has optimal activity outside of the normal physiological range of pH and temperature, we wanted to expand upon these findings. In [Figure 8](#), we show that GST-HsPRMT7 activity decreases not only with NaCl but also with KCl. To ask if this effect is due to ionic strength, we also assayed the enzyme in Na_2SO_4 . We found that the activity is dependent on the ionic strength with half activity at about 50 mM and almost no activity at 150 mM. Since typical intracellular ionic strength values range from about 130 to 270 mM (14), these results

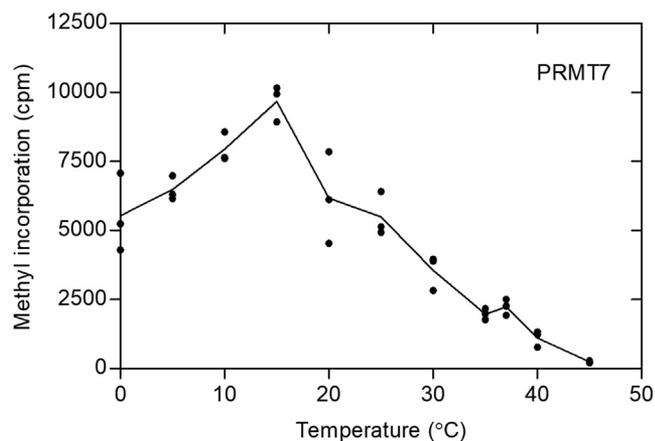


Figure 3. Human PRMT7 is more active at lower temperatures under steady-state kinetics. P81-based assay of GST-HsPRMT7, 10 μ M HsH2B (23–37), and 0.14 μ M [3 H]AdoMet was carried out as described in [Experimental procedures](#). Reaction mixtures were incubated for 30 min, spotted on P81 phosphocellulose paper, and further assayed as described in [Experimental procedures](#). The data shown are from a single experiment; similar results were seen in three additional replicated experiments. HsH2B, histone H2B; PRMT, protein arginine methyltransferase.

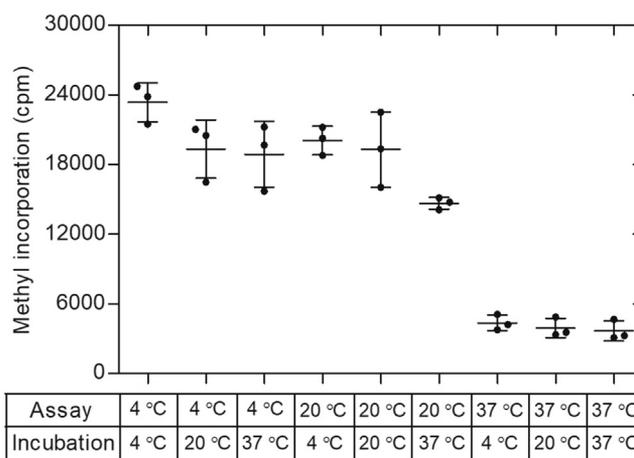


Figure 4. Human PRMT7 activity is stable from 4 $^{\circ}$ C to 37 $^{\circ}$ C. GST-HsPRMT7 was preincubated for 2 h at 4 $^{\circ}$ C, 20 $^{\circ}$ C, and 37 $^{\circ}$ C in the presence of the HsH2B (23–37) substrate and 50 mM K-Hepes, 1 mM DTT at a final pH of 8.5. The enzymatic reaction was then initiated by addition of a final concentration of 0.7 μ M [3 H]AdoMet and was then incubated for 2 h at the indicated temperature followed by the P81 phosphocellulose assay as described in [Experimental procedures](#). Data are shown for triplicate of samples with standard deviation shown as error bars. HsH2B, histone H2B; PRMT, protein arginine methyltransferase.

Stress and protein arginine methyltransferase enzymology

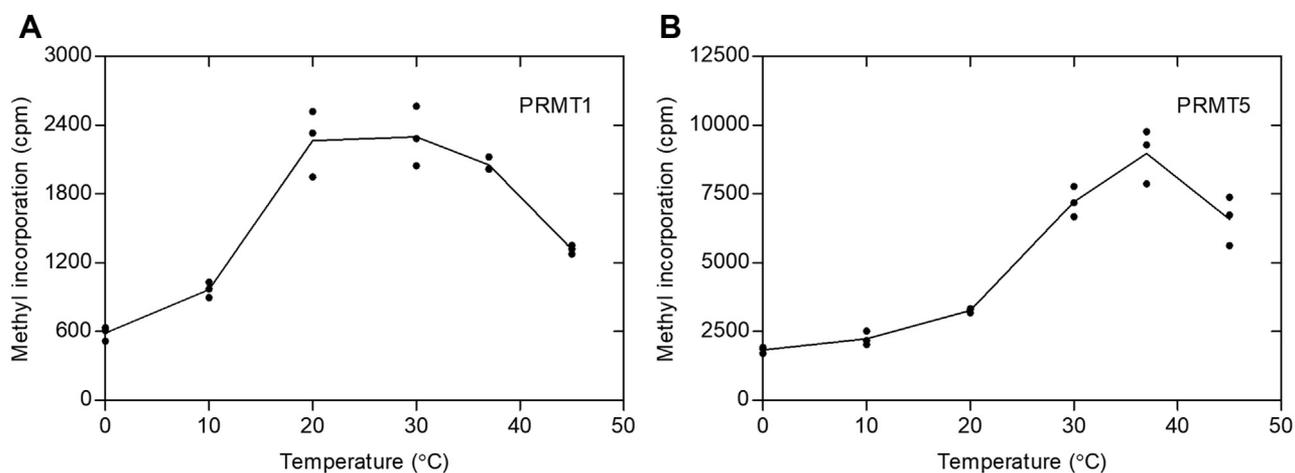


Figure 5. Human PRMT1 and PRMT5/MEP50 are more active near physiological temperatures. *In vitro* reaction mixtures were incubated at the specified temperatures and assayed using a P81 phosphocellulose assay. Reaction mixtures contained 50 mM K-HEPES, 1 mM DTT and had a final pH of 8.0. Reactions were allowed to incubate for 30 min. **A**, 80 nM of His-HsPRMT1 was incubated with 10 μ M HsH4 (1–21) and 0.14 μ M [3 H]AdoMet. **B**, 5.6 nM of HsPRMT5/MEP50 was incubated with 10 μ M of HsH4 (1–21), and 0.14 μ M [3 H]AdoMet. HsH4, histone H4; PRMT, protein arginine methyltransferase.

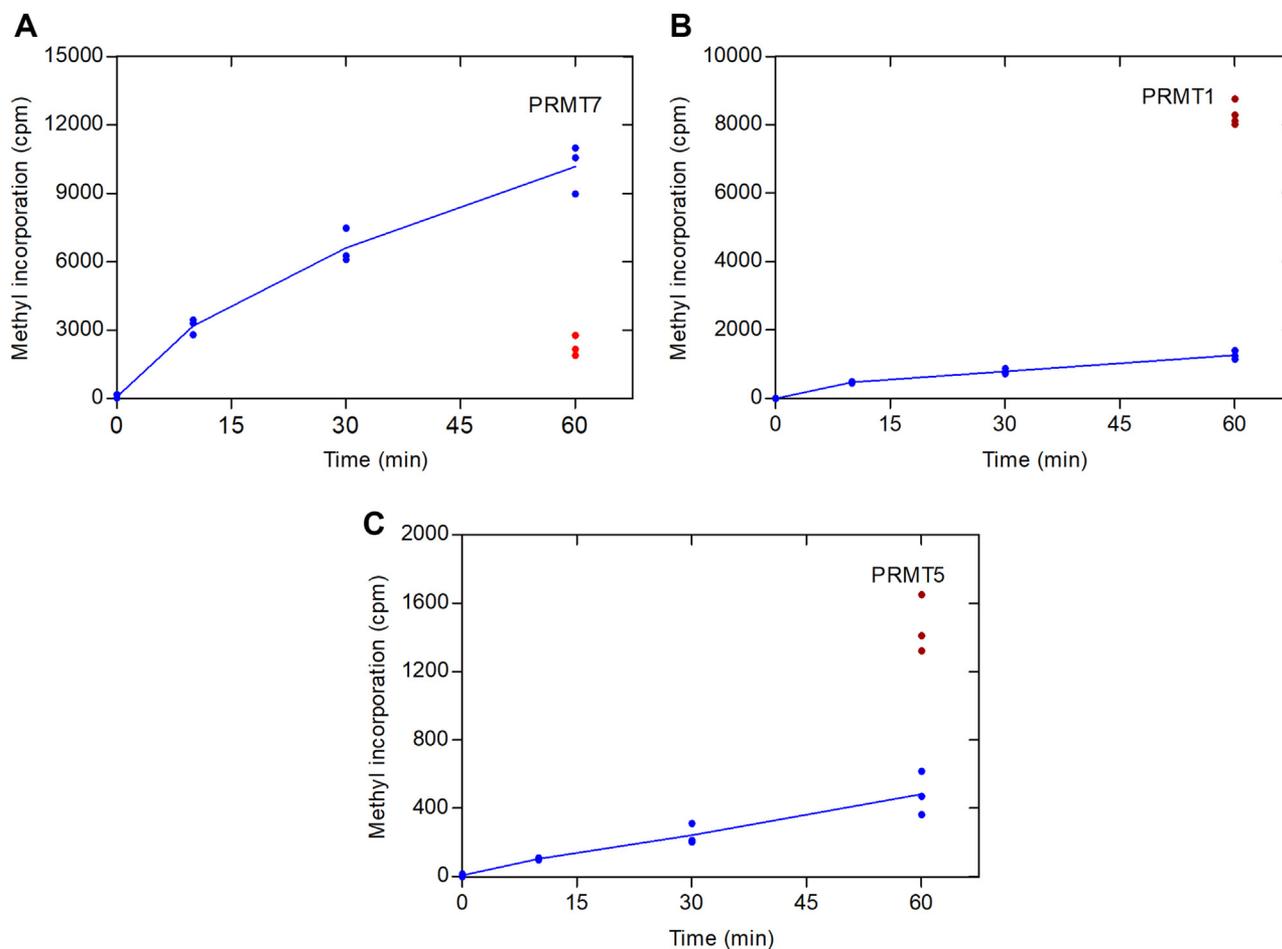


Figure 6. Human PRMT7, PRMT1, and PRMT5/MEP50 are active at 0 °C. P81 phosphocellulose assays of reactions containing GST-HsPRMT7 with HsH2B (23–37), His-HsPRMT1 with HsH4 (1–21), or HsPRMT5/MEP50 with HsH4(1–21) was carried out as described in [Experimental procedures](#). **A**, Five micrograms of GST-HsPRMT7 was mixed with 10 μ M HsH2B (23–37) and 0.14 μ M of [3 H]AdoMet. Reaction mixtures contained 50 mM K-Hepes, 1 mM DTT, at a final reaction pH of 8.5. **B**, 40 nM of His-HsPRMT1 was mixed with 10 μ M HsH4 (1–21) and 0.14 μ M of [3 H]AdoMet. Reaction mixtures contained 50 mM K-Hepes, 1 mM DTT, at a final reaction pH of 7.0. **C**, 5.6 nM of HsPRMT5/MEP50 was mixed with 10 μ M HsH4 (1–21) and 0.14 μ M [3 H]AdoMet. Reaction mixtures contained 50 mM K-Hepes, 1 mM DTT, at a final reaction pH of 7.0. All reaction mixtures were equilibrated to 0 °C for 2 min prior to initiation with [3 H]AdoMet. The *blue line* represents averages of replicates at 0 °C. *Red dots* represent reactions incubated at 37 °C. HsH2B, histone H2B; HsH4, histone H4; PRMT, protein arginine methyltransferase.

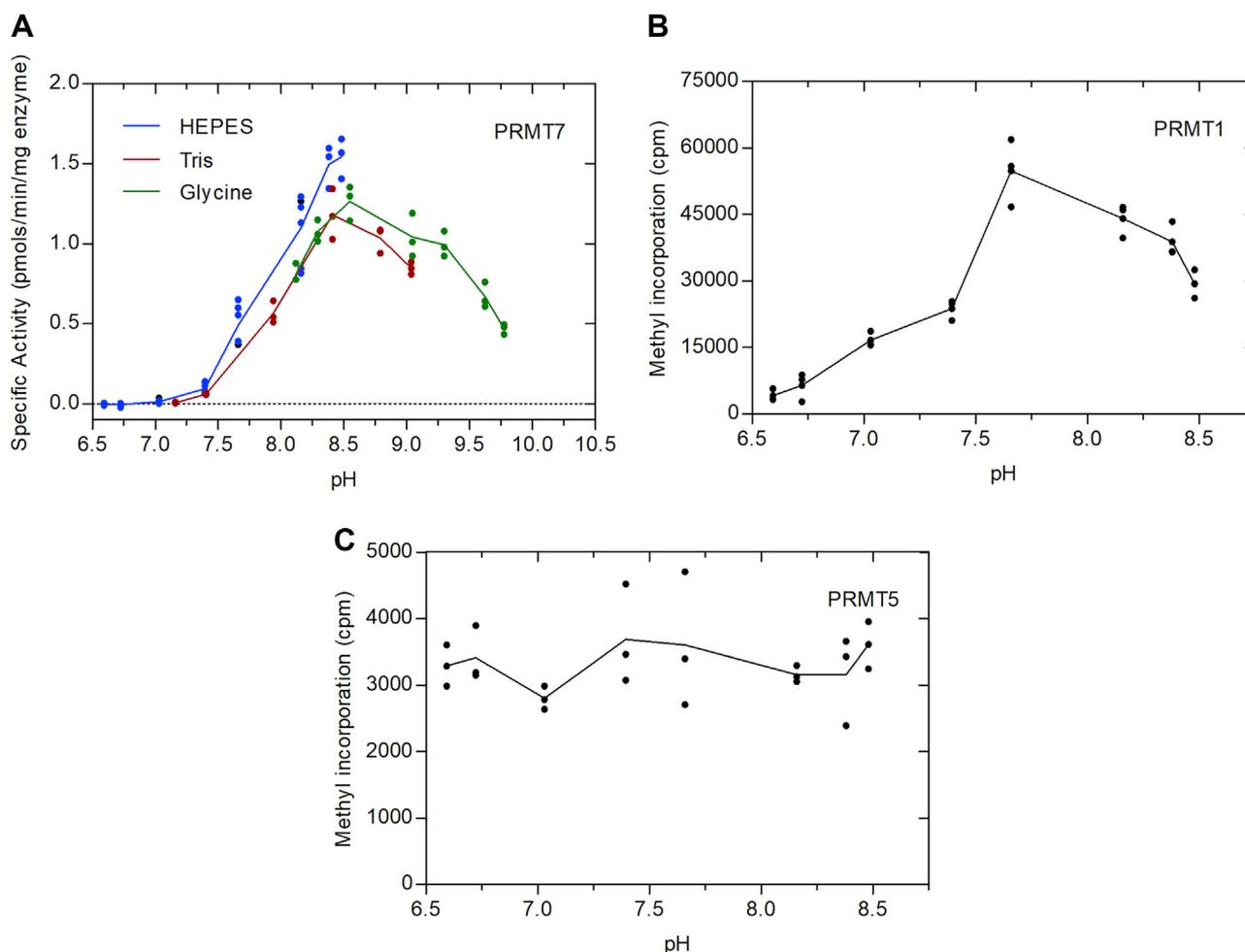


Figure 7. pH dependence of human PRMT7, PRMT1, and PRMT5/MEP50. P81 phosphocellulose assay with either GST-HsPRMT7 and HsH2B (23–37), His-HsPRMT1 and HsH4 (1–21) or HsPRMT5/MEP50 and HsH4 (1–21) was carried out as described in [Experimental procedures](#). **A**, five micrograms of GST-HsPRMT7, 10 μ M of HsH2B (23–37), and 0.14 μ M [3 H]AdoMet. The *red line* represents averages of replicates where reaction mixtures contain 50 mM K-Hepes, 1 mM DTT. The *green line* represents averages of replicates reaction mixtures containing 50 mM glycine and 1 mM DTT. The *blue line* represents averages of replicates where reaction mixtures contain 50 mM Tris and 1 mM DTT. Reaction mixtures were incubated at 20 $^{\circ}$ C for 60 min. **B**, 40 nM of His-HsPRMT1, 10 μ M HsH4 (1–21), and 0.14 μ M [3 H]AdoMet were mixed with 50 mM K-Hepes and 1 mM DTT and incubated at 37 $^{\circ}$ C for 30 min. **C**, 5.6 nM of HsPRMT5/MEP50, 10 μ M of HsH4 (1–21) and 0.14 μ M [3 H]AdoMet were mixed with 50 mM K-Hepes and 1 mM DTT and incubated at 37 $^{\circ}$ C for 30 min. The *black line* represents the averages of *black dot* replicates for His-HsPRMT1 and HsPRMT5/MEP50. In all cases, the final pH was determined from mock mixtures of buffer components, the buffer used in the enzyme preparation, and the solvent for the [3 H]AdoMet. HsH2B, histone H2B; HsH4, human histone H4; PRMT, protein arginine methyltransferase.

suggest that PRMT7 activity is generally repressed under physiological conditions. Control experiments showed that changes in the ionic strength of the incubation mixture did not affect the efficiency of the P81 phosphocellulose assay ([Fig. S3](#)). We also performed assays on the effect of ionic strength using GST-GAR as a substrate and the SDS-PAGE assay ([Fig. S4](#)). We note that the loss of activity with ionic strength is less with the GST-GAR substrate than with the HsH2B (23–37) peptide, confirming the results of [Feng *et al.* \(16\)](#).

We then examined the effect of calcium and magnesium ions on the activity of PRMT7. Here we found inhibition greater than expected from the ionic strength. With calcium ions, we found half inhibition at approximately 0.7 mM, and with magnesium ions, we found half inhibition at about 1.6 mM ([Fig. 8](#)). Since intracellular calcium ion concentrations are submicromolar, we might expect little effect on PRMT activity except perhaps when cells undergo intracellular

calcium release. Intracellular free magnesium concentration has been estimated in the cytosol of mammalian cells at about 0.5 mM ([19](#)), suggesting little inhibition of PRMT7 with changes in magnesium ion concentration. Taking all of these results together, it appears that PRMT7 may only be active under conditions where intracellular temperature, pH, and ionic conditions are disrupted.

PRMT1 and PRMT5 are also inhibited by the addition of salts

Finally, we sought to determine whether PRMT1 and PRMT5 activity was also affected by ionic strength or calcium and magnesium ions ([Fig. 9](#)). For His-HsPRMT1 with the human histone H4 (1–21) peptide, we found half maximal inhibition by ionic strength at about 100 mM and half maximal inhibition by calcium and magnesium ions at about 2 mM. For HsPRMT5/MEP50 under similar conditions, we found half

Stress and protein arginine methyltransferase enzymology

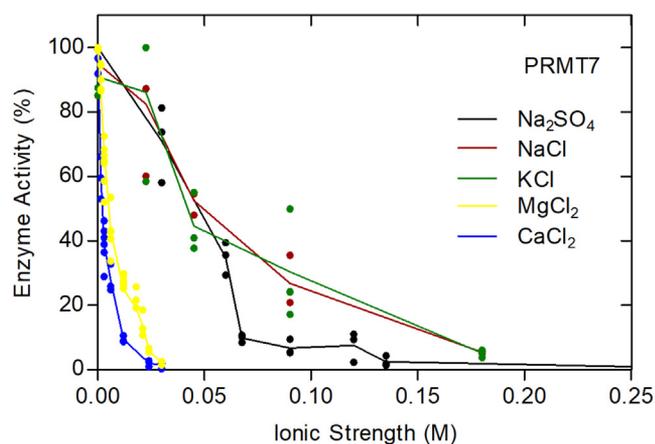


Figure 8. Human PRMT7 activity decreases with ionic strength. Five micrograms of GST-HsPRMT7 was mixed with 10 μM HsH2B (23–37) and 0.7 μM [^3H]AdoMet. All reaction mixtures contained 50 mM K-Hepes and 1 mM DTT with a final pH of 8.5. The *black line* represents averages when Na_2SO_4 was added to reaction mixtures. The *red line* represents averages when NaCl was added to reaction mixtures. The *green line* represents averages when KCl was added to reaction mixtures. The *yellow line* represents averages when MgCl_2 was added to reaction mixtures. The *blue line* represents the averages when CaCl_2 was added to the reaction mixtures. All reactions were incubated for 10 min at 20 $^\circ\text{C}$. HsH2B, histone H2B; PRMT, protein arginine methyltransferase.

maximal inhibition by ionic strength at about 120 mM, half maximal inhibition of calcium ions at about 5 mM, and half maximal inhibition of magnesium ions at about 8 mM. Interestingly, these enzymes are also inhibited in the physiological ionic strength range, although the inhibition by calcium and magnesium ions is less than that seen with PRMT7. It appears that PRMT1 and PRMT5 would not be significantly inhibited by physiological levels of calcium and magnesium.

Discussion

In this study, we were surprised to find that human PRMT1, PRMT5, and PRMT7 had optimal catalytic activities *in vitro*

under nonphysiological conditions. For example, all three enzymes displayed markedly reduced activity at ionic strengths comparable to the cytosol of mammalian cells. While PRMT1 and PRMT5 were active at a range of pH values, PRMT7 had little activity at the expected intracellular pH of 7.2 and was only fully active at alkaline pH values. Finally, all of the three enzymes displayed a significant catalytic activity at low temperatures, including 0 $^\circ\text{C}$. Significantly, human PRMT7 had markedly reduced activity at the physiological temperature of 37 $^\circ\text{C}$, while PRMT1, PRMT5, and PRMT9 (17) had near-optimal activity at 37 $^\circ\text{C}$. We note that the PRMT7 from the soil nematode worm *Caenorhabditis elegans* has a distinct temperature profile with optimal activity at about 20 $^\circ\text{C}$ near that of its environment in nature (17).

We wanted to know if the optimal activity of PRMTs under these nonphysiological conditions would mean that PRMTs are responding to environmental stresses. We wanted to ask under what conditions would PRMTs be exposed to temperatures lower than 37 $^\circ\text{C}$, under what conditions would PRMTs be exposed to pH values above 7.2, and under what conditions would PRMTs be exposed to ionic strengths lower than what is generally found in mammalian cells.

Humans generally maintain their temperature at 37 $^\circ\text{C}$ even when exposed to environmental cold. However, skin cells and cells lining the lungs can be significantly affected (20, 21). We note that mammalian cells have “cold shock” proteins that are induced by a cold shock response (22–24). One example of cold response is that optimal human sperm production is maintained when the testis is 2 to 7 $^\circ\text{C}$ below body temperature (25). It has been suggested that the TRPM8 sensor protein is important for the cold shock response in testes (21). Moderately reduced temperatures have also been found to suppress apoptosis of neural stem cells although it is not clear when such cells would be exposed to lower temperatures (26).

Interestingly, a comparison of genes has associated PRMT7, along with a small group of other genes, to the adaptation of

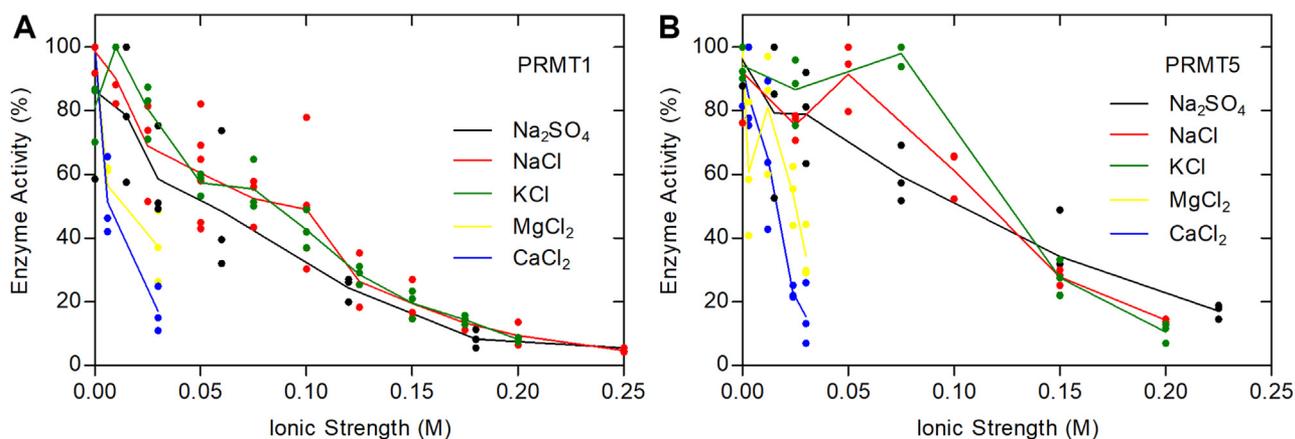


Figure 9. Human PRMT1 and PRMT5/MEP50 activities are reduced with increasing ionic strength. P81 phosphocellulose of His-HsPRMT1 or HsPRMT5/MEP50 with HsH4 (1–21) was carried out as described in [Experimental procedures](#). A, 40 nM of His-HsPRMT1, 10 μM HsH4 (1–21), and 0.7 μM [^3H]AdoMet were incubated with 50 mM K-Hepes and 1 mM DTT at a final pH of 7.5. Reaction mixtures were incubated at 37 $^\circ\text{C}$ for 15 min. B, 5.6 nM of HsPRMT5/MEP50, 10 μM HsH4 (1–21), and 0.7 μM [^3H]AdoMet were incubated with 50 mM K-Hepes and 1 mM DTT at a final pH of 7.5. Reaction mixtures were incubated at 37 $^\circ\text{C}$ for 30 min. The *black line* represents averages when Na_2SO_4 was added to reaction mixtures. The *red line* represents averages when NaCl was added to reaction mixtures. The *green line* represents averages when KCl was added to reaction mixtures. The *yellow line* represents averages when MgCl_2 was added to reaction mixtures. The *blue line* represents the averages when CaCl_2 was added to the reaction mixtures. HsH4, human histone H4; PRMT, protein arginine methyltransferase.

elephantid species to cold environments, particularly of mammoths to the arctic (27).

Many mammalian proteins have been shown to be denatured at subphysiological temperatures (28). However, it is not uncommon for enzymes to adapt to low temperatures, especially those in fish and in bacteria exposed to cold environments (29).

To determine how common is cold activation of enzymes, we searched the literature for publications reporting on the temperature dependence of human or mammalian S-adenosylmethionine methyltransferases. While many of these enzymes show optimal activities around 37 °C as expected, we were interested to find an example of an enzyme that displayed cold activation similar to what we report here for PRMT7. This enzyme is the NS5 RNA methyltransferase from the Dengue virus 4. Optimal activity was found between 22 °C and 30 °C, with less than 20% of this activity found at 37 °C (30). It is presently unclear what the biological advantage is for PRMT7 or the RNA methyltransferase to be cold activated. Additionally, it is unknown why PRMT1, PRMT5, and PRMT7 have significant enzymatic activity at 0 °C.

PRMT7 was shown here to have very little activity at the expected cytosolic pH of about 7.2 in mammalian cells (13, 31). The activity of PRMT7 at pH 7.2 was found to be less than 5% of its optimal activity at pH 8.4. In comparison to pH 7.2, human PRMT1 has about 30% of its optimal activity at pH 7.6, while PRMT5 has a broad optimal range between 6.5 and 8.5. Interestingly, the Dengue virus 4 NS5 RNA methyltransferase also shows enhanced activity at alkaline pH. Its activity at 7.2 is only about 20% of its optimal activity at pH 9.0 (30). A similar situation occurs with the PRDM9 histone H3 lysine 36 methyltransferase where the activity at pH 7.2 is only about 15% of the optimal activity at pH 8.5 (32). All of these results suggest that PRMT7, the NS5 RNA methyltransferase, the histone lysine methyltransferase, and possibly PRMT1 may be regulated by conditions resulting in the alkalization of cells.

PRMT1, PRMT5, and PRMT7 all show inhibition with increasing ionic strength. For PRMT7, this inhibition was much more pronounced with the HsH2B (23–37) peptide than for the GST-GAR substrate. We were intrigued to find that Dengue virus-4 RNA methyltransferase and the PRDM9 histone lysine methyltransferase also showed sensitivity to ionic strength (30, 32). The physiological relevance of such sensitivity is not clear although it has been suggested that cells can respond to changes in ionic strength (14). Normal intracellular ionic strength has been estimated at about 110 to 130 mM in the human embryonic kidney cell line HEK293 reflecting the known intracellular sodium ion and potassium ion levels (14). At this ionic strength, PRMT7 (at least with the HsH2B (23–37) peptide substrate) and the histone lysine methyltransferase PRDM9 have less than 10% activity, while PRMT1, PRMT5, and the Dengue-4 RNA methyltransferase have less than 50% activity.

Finally, PRMT1, PRMT5, and PRMT7 are all inhibited by increasing concentrations of magnesium ions. PRMT7 was found to be inhibited by magnesium ion concentration with half activity found at 1.6 mM, while PRMT1 was half inhibited

at 2 mM and PRMT5 was half inhibited at 8 mM. Since the intracellular free magnesium ion concentration has been estimated at about 0.5 mM (19), it appears that the inhibition seen here would not be physiologically relevant. These methyltransferases are also inhibited by calcium ions at levels much greater than the levels found intracellularly.

At present, we do not understand the importance of the subphysiological activities of PRMT1, PRMT5, and PRMT7 and the RNA and protein lysine methyltransferases as described earlier in the study. The similarities in the responses to temperature, pH, and ionic strength are, however, remarkable.

Evidence has been presented that physiological levels of hydrogen peroxide can inhibit human PRMT1 and that this inhibition can be reversed by reducing agents, suggesting a possible regulation of PRMT1 by redox reactions (33). It is clear that PRMT7 activity is dependent upon a reducing environment as well (17, 34). In all of our experiments, we performed incubations in 1 mM DTT (17).

It remains to be shown whether or not PRMTs, especially PRMT7, are regulated *in vivo* by intracellular temperature, pH, or ionic strength. Unfortunately, it has been challenging to measure the individual activities of the nine family members of mammalian PRMTs in cells cultured under normal or low temperature conditions, or under conditions that would affect their intracellular ionic strength or pH. This is particularly a problem for PRMT7, whose activity appears to be generally low in comparison with the major type I PRMT1 enzyme or the major type II PRMT5 enzyme (5, 35). Although evidence has been presented for a number of *in vivo* sites of PRMT7 methylation (7, 10, 36–38), it is not clear so far that these are specifically modified by PRMT7 alone and are not also sites for methylation by other PRMTs (39). Additionally, PRMT7 methylation of Arg-17 of a histone H4 peptide has been shown to allosterically activate PRMT5 methylation at the distinct Arg-3 site (9). Such cross talk between PRMTs can complicate the analysis of specific PRMT activity in cells. Finally, it is now clear that reductions in the activity of one PRMT can result in the methylation of its sites by other members of the PRMT family (40, 41). The identification of specific substrates and the generation of specific antibodies raised against their monomethylation site(s) will be important steps in quantitating PRMT7 activity in cells and testing how this activity may respond to changes in their intracellular environment.

Experimental procedures

Preparation of enzymes and protein substrates

Recombinant *H. sapiens* PRMT7 DNA was cloned into a pGEX-2T plasmid and expressed in BL21 DE3 *E. coli*. This plasmid encodes the full sequence (residues 1–218) of glutathione S-transferase from *Schistosoma japonicum* followed by the linker SDLVPRGSST and the full sequence (residues 1–692) of human PRMT7 and was designated as GST-HsPRMT7. This plasmid is available from Addgene.org #34693. Cells were grown in 2L of LB media at 37 °C to an A_{600} nm of 0.6 to 0.8 in the presence of 100 µg/ml ampicillin.

Stress and protein arginine methyltransferase enzymology

Isopropyl-beta-D-thiogalactoside (CAS # 367–93–1, GoldBio) was then added to a final concentration of 1 mM, and the culture was incubated for an additional 16 to 20 h at 20 °C. Cells were harvested by centrifugation, and 0.5 g of cells was resuspended in 40 ml of 140 mM NaCl, 2.7 mM KCl, 10 mM Na₂HPO₄, and 1.8 mM KH₂PO₄, pH = 7.5, supplemented with an EDTA-free protease inhibitor tablet (Catalog No. PIA32965, Thermo Scientific). The cell suspension was lysed using a Fisher Scientific 550 Sonic Dismembrator with a microtip attachment in four cycles on ice for 1 min, 0.5 s on pulse, and 0.5 s off pulse at 25% amplitude. The lysate was then centrifuged at 27000g for 1 h, and the supernatant was loaded onto a manually packed column (ID 1.5 cm × 15 cm, 4 ml bed volume) containing glutathione Sepharose 4B resin (Catalog No. 17075601, GE Healthcare Life Sciences) equilibrated in the aforementioned buffer at 22 °C. Enzyme was eluted with 10 mM glutathione and 50 mM Tris-HCl, pH 8.0. Purified GST-HsPRMT7 was then dialyzed overnight in 50 mM K-Hepes, 1 mM DTT, pH 8.0, at 4 °C. GST-HsPRMT7 was then mixed with glycerol to a final concentration of 20% and stored at –80 °C. The total GST-HsPRMT7 concentration was determined using a NanoDrop 2000c spectrophotometer by the absorption at 280 nm, assuming that 1 mg/ml would give 1 A.

An N-terminal His-tagged construct of human PRMT1 (designated His-HsPRMT1) was prepared as described previously and purified using a HisTrap Fast Flow (Cytiva 17–5255–01) as per the manufacturer's protocol (42).

Human PRMT5/MEP50 (HsPRMT5/MEP50) FLAG-tag His-tag complex protein was purchased from BioSciences as recombinantly coexpressed and purified proteins in HEK293T cells (0.64 mg/ml, Catalog No. 51045, Lot 170407A; BPS Biosciences), formulated in 40 mM Tris-HCl, pH 8.0, 110 mM NaCl, 2.2 mM KCl, 80 µg/ml FLAG peptide, 20% glycerol, and 3 mM DTT.

A fusion protein of glutathione S-transferase followed by the linker SDLVPRGS and human fibrillarin residues 1 to 145 with substitutions K2E and A145V (GST-GAR) was prepared by expression of plasmid #34697 of [Addgene.org](https://addgene.org/17/). (17). The protein was purified as described for GST-HsPRMT7 above.

Human HsH2B was purchased from NEB, Cat. No. M2505S, as a recombinant protein expressed in *E. coli*, stored in 300 mM NaCl, 1 mM EDTA, and 20 mM sodium phosphate at pH 7.

Peptides

Synthetic peptides HsH2B (23–37) (Ac-KKDGKKRKR SRKESY, 98.6% purity, MW: 1936.23, Lot No. P17171304) and human histone H4 (1–21) (Ac-SGRGKGGKGLGKGGAKR HRKV, 96.2% purity, MW: 2133.47, Lot No. 89488400001/PE1586) were purchased from Genescript.

In vitro methylation reactions using P81 phosphocellulose

GST-HsPRMT7, His-HsPRMT1, or PRMT5/MEP50 was mixed with a peptide derived from residues 23 to 37 of HsH2B, GST-GAR, or a peptide derived from the acetylated N-

terminus of histone H4 (residues 1–21). Reactions were initiated with *S*-adenosyl-L-[methyl-³H]methionine ([³H]AdoMet) (81.9 Ci/mmol, 7 µM in 9:1 10 mM H₂SO₄:ethanol; Catalog No. NET155H001MC, PerkinElmer) in a final volume of 30 µl. The reaction mixtures were equilibrated to the desired temperature for 2 min prior to the addition of [³H]AdoMet and incubation in a temperature-regulated water bath. Post incubation, the reaction mixtures were terminated with 0.5 µl of 100% trifluoroacetic acid and spun down for 10 s at 10,600g in an Eppendorf microcentrifuge (Model 5415C). All reaction mixtures contained 1 mM DTT and unless otherwise described a final concentration of 50 mM K-HEPES at a final pH of 8.5. To determine the final pH of reaction mixtures, mock mixtures were created containing the appropriate volumes of the enzyme buffer (50 mM K-Hepes, 1 mM DTT, pH 8.0), the [³H]AdoMet solvent (18 mM HCl), and the reaction buffer containing 1 mM DTT and 50 mM K-Hepes, 50 mM glycine, or 50 mM Tris-HCl.

The extent of methylation was determined by a P81 phosphocellulose assay. Twenty-five microliters of reaction mixture was spotted on to a 1 × 1-cm² of phosphocellulose P81 paper (Catalog No. 05–717-2A, Lab Alley). Papers were dried for 90 min at room temperature and were then washed 3 times with 300 ml of 50 mM sodium bicarbonate, pH = 9.0. The papers were dried again for 90 min in 20-ml scintillation vials (Catalog No. 66021-704, VWR), and 5 ml of scintillation fluid (111177-CS, RPI) was added. Vials were counted in a Beckman LS 6500 Liquid Scintillation Counter for four cycles at 5 min per cycle. The sum of enzyme-only and substrate-only controls was subtracted from individual replicates.

In vitro methylation gel assay

Recombinant GST-HsPRMT7 was incubated with either GST-GAR as described earlier in the study or recombinant human HsH2B expressed in *E. coli* (Catalog No. M2505S, New England BioLabs) in a total volume of 30 µl. Post incubation, the reaction mixtures were spun down for 10 s at 10,600g. The reactions were then terminated with 5 µl of 5× sample buffer: 250 mM Tris-HCl, 10% SDS, 30% glycerol, 500 mM DTT, and 0.05% bromophenol blue and heated for 1 min in a 95 °C sand bath. The reaction mixtures were allowed to cool to room temperature before loading either onto a 4 to 12% 15-well ExpressPlus PAGE gel (Catalog No. NC1486979, Fisher Scientific Co) or 4 to 20% 10-well Express PAGE gel (Catalog No. NC0613877, Fisher Scientific Co). The gel was allowed to run for 1 h at 140 V, 400 mA using a BioRad PowerPac 300 electrophoresis Power supply. The gels were stained for 1 h in Coomassie stain: 50% methanol and 10% acetic acid, followed by an overnight destain consisting of 15% methanol and 10% acetic acid. The gels were then soaked in water for minimum 3 h prior to enhancing with En³Hance (Catalog No. 6NE9701, PerkinElmer) for 30 min. The gels were then put into water for 1 h post enhancement, wrapped in gel-drying film (Catalog No. PR-V713, Promega), and dried in a Model 583 Gel Dryer, Biorad for 2 h at 80 °C. The dried gel was placed in an autoradiography cassette, and an HyBlot CL autoradiography film

(Catalog No. NC9556985, Thermo Scientific) was placed on top of the gel. The film was exposed for 3 to 30 days at -80°C before development

Data availability

All data described in the manuscript are contained within the manuscript. Additional data are available upon request.

Supporting information—This article contains supporting information.

Author contributions—T. L. L. and S. G. C. conceptualization; T. L. L. and S. G. C. methodology; T. L. L. and S. G. C. formal analysis; T. L. L. investigation; T. L. L. resources; T. L. L. and S. G. C. writing - original draft; T. L. L. and S. G. C. writing - review & editing; T. L. L. and S. G. C. visualization; S. G. C. supervision; S. G. C. project administration; S. G. C. funding acquisition.

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Conflict of interest—The authors declare that they have no conflicts of interest with the content of this article.

Abbreviations—The abbreviations used are: HsH2B, human histone H2B; PRMT, protein arginine methyltransferase.

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Stress and protein arginine methyltransferase enzymology

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Supporting information

Human Protein Arginine Methyltransferases (PRMTs) Can Be Optimally Active Under Non-Physiological Conditions

Troy L. Lowe and Steven G. Clarke

List of material included:

Figure 1S: pH Does not affect the P81 methylation assay efficiency.

Figure 2S: pH dependence of human PRMT7 with a protein substrate.

Figure 3S: Salts do not affect the P81 methylation assay efficiency.

Figure 4S: Human PRMT7 activity decreases with ionic strength with a GST-GAR substrate.

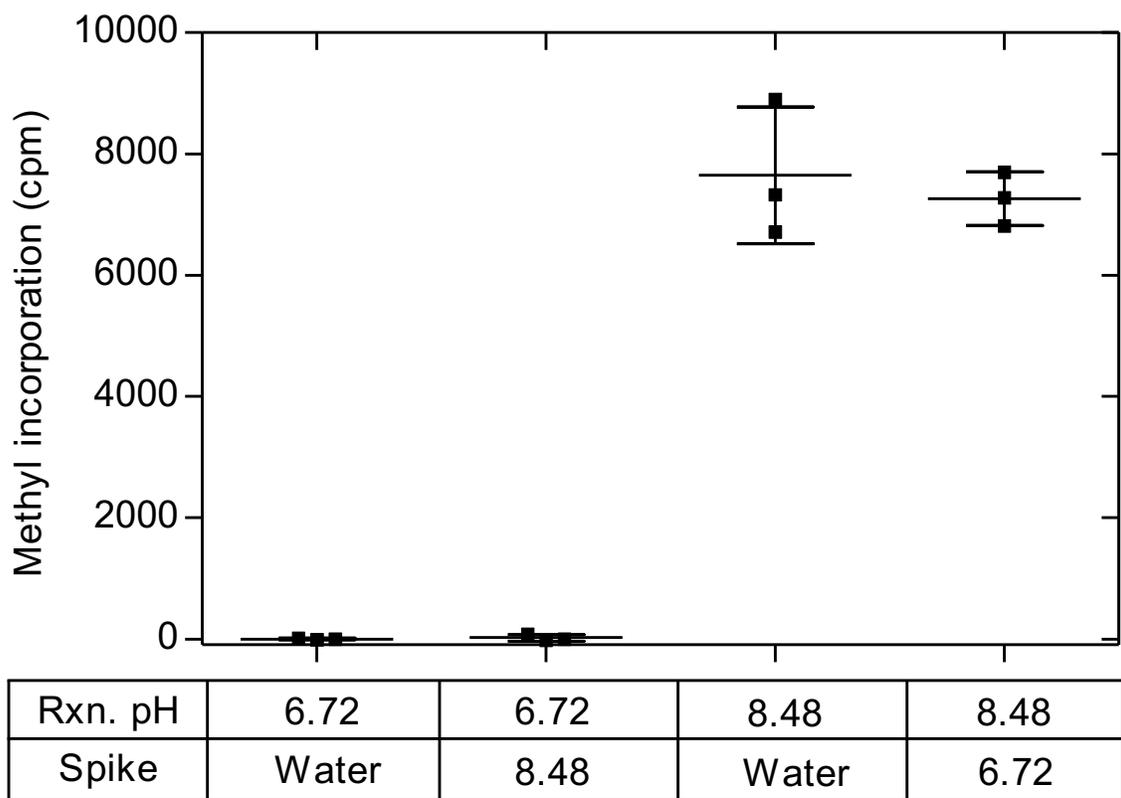


Figure 1S: pH does not affect the P81 methylation assay efficiency. Replica reaction mixtures in a final volume of 30 μ L were prepared containing 5 μ g GST-HsPRMT7, 10 μ M HsH2B (23-37), 50 mM K-HEPES, 1 mM DTT, and 0.7 μ M [3 H]AdoMet at a final pH of 8.48, or a final pH of 6.72. These reaction mixtures were incubated for 1 hr at 20 $^{\circ}$ C and were then terminated by the addition of 0.5 μ L of 100% TFA. At this point, either 3 μ L of water or 3 μ L of 500 mM K-HEPES, pH 8.48 or 6.72 were added and the assay performed as described in the “Experimental Procedures” section. ANOVA analysis showed no significant difference between the activity with the addition of water or buffer with a pH of 8.48 or 6.72.

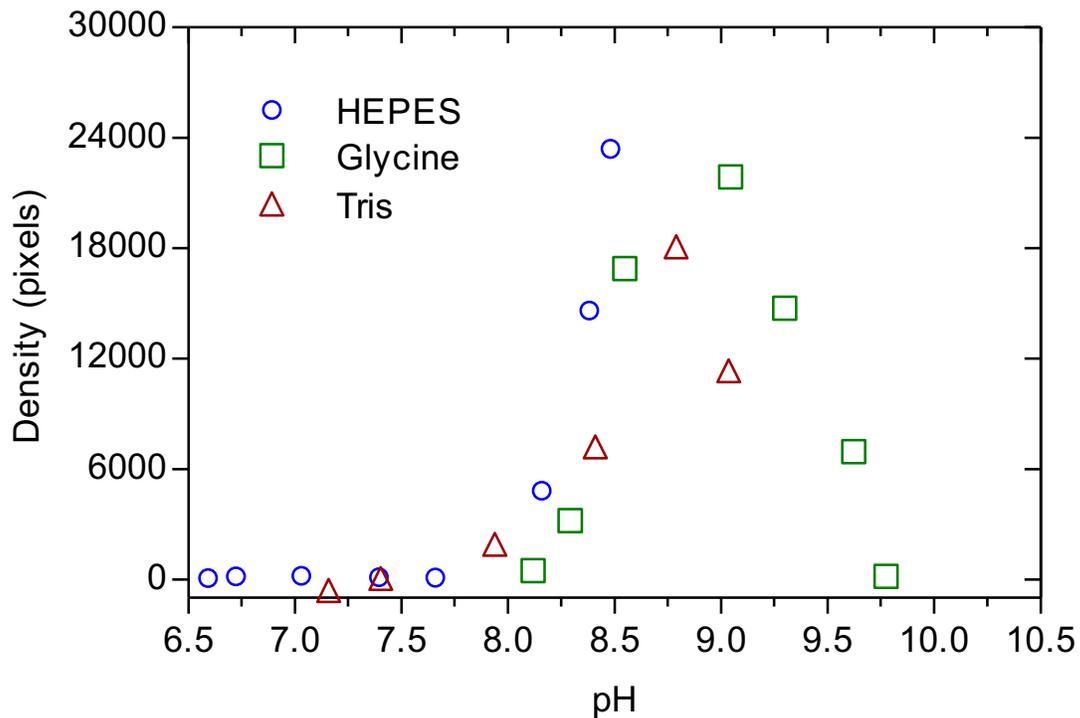


Figure 2S: pH dependence of human PRMT7 with a protein substrate. GST-HsPRMT7 5 μg was incubated with 6.2 μg GST-GAR, 1 mM DTT, 50 mM of the indicated buffer, and 0.14 μM [^3H]AdoMet in a final volume of 30 μL . The reaction was performed for 1 h at 20 $^{\circ}\text{C}$. The reaction was then quenched by the addition of SDS sample buffer and the mixtures were fractionated by SDS-PAGE as described in “Experimental Procedures.” Films were analyzed by densitometry using ImageJ software after scanning. Blue open circles represent samples incubated in K-HEPES at the indicated final pH values, while red open triangles represent samples incubated in Tris-HCl, and green open squares represents samples incubated in a glycine buffer.

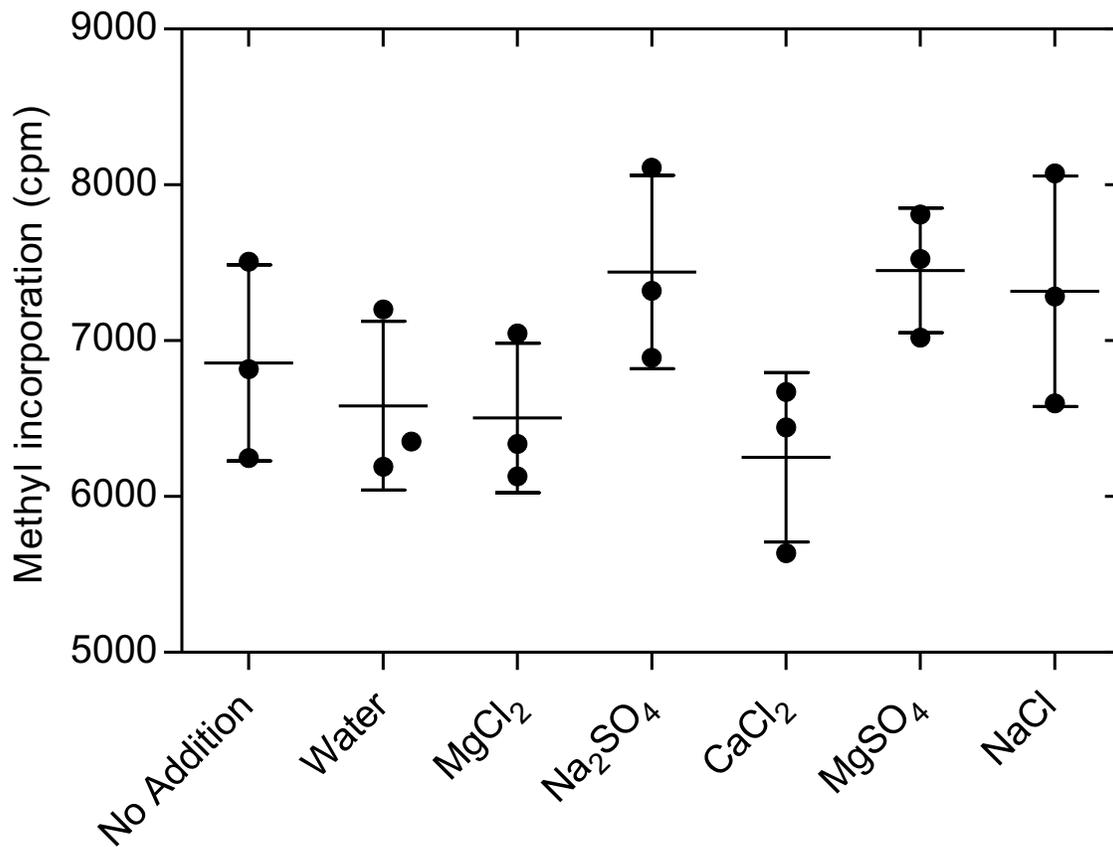


Figure 3S: Salts do not affect the P81 methylation assay efficiency. Replica reaction mixtures in a final volume of 30 μ L were prepared containing 5 μ g GST-HsPRMT7, 10 μ M HsH2B (23-37), 50 mM K-HEPES, 1 mM DTT, and 0.7 μ M [³H]AdoMet at a final pH of 8.5. These reaction mixtures were incubated for 1 hr at 25 °C and were then terminated by the addition of 0.5 μ L of 100% TFA. At this point, either no addition was made or 3 μ L of water, 3 μ L of 300 mM MgCl₂, Na₂SO₄, or CaCl₂, or 3 μ L of 900 mM NaCl, or 3 μ L of 225 mM MgSO₄ were added and the assay performed as described in the “Experimental Procedures” section. ANOVA analysis showed no significant difference between the activity with no addition or with the water or salt additions.

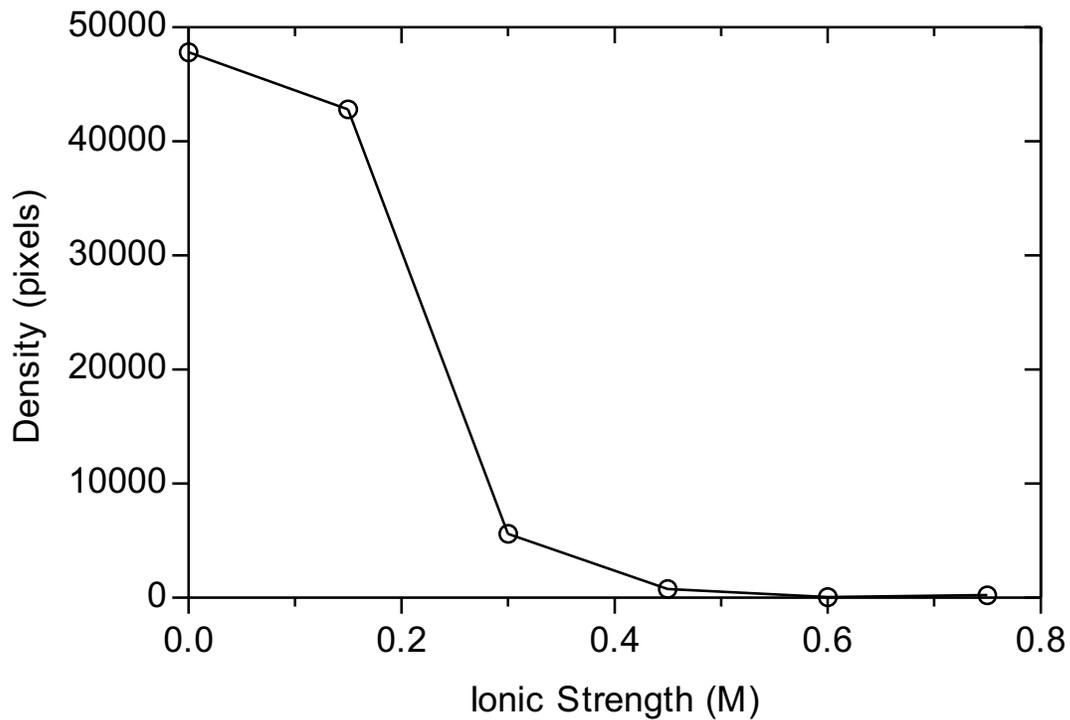


Figure 4S: Human PRMT7 activity decreases with ionic strength with a GST-GAR substrate. 5 μg GST-HsPRMT7 was incubated with 6.2 μg GST-GAR, 1 mM DTT, 50 mM K-HEPES, and 0.14 μM [^3H]AdoMet in a final volume of 30 μL and a final pH of 8.5. The ionic strength was adjusted with sodium sulfate to the indicated level. The reaction was performed for 1 h at 20 $^{\circ}\text{C}$. The reaction was quenched by the addition of SDS sample buffer and the mixtures were fractionated by SDS-PAGE as described in “Experimental Procedures.” Films were analyzed by densitometry of the GST-GAR using ImageJ software after scanning.