

Design, Synthesis, and Antiviral Activity of 2'-Deoxy-2'-fluoro-2'-C-methylcytidine, a Potent Inhibitor of Hepatitis C Virus Replication

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The pyrimidine nucleoside beta-D-2'-deoxy-2'-fluoro-2'-C-methylcytidine (**1**) was designed as a hepatitis C virus RNA-dependent RNA polymerase (HCV RdRp) inhibitor. The title compound was obtained by a DAST fluorination of *N*⁴-benzoyl-1-(2-methyl-3,5-di-*O*-benzoyl-β-D-arabinofuranosyl)cytosine (**6**) to provide *N*⁴-benzoyl-1-[2-fluoro-2-methyl-3,5-di-*O*-benzoyl-β-D-ribofuranosyl]cytosine (**7a**). The protected 2'-C-methylcytidine (**7c**) was obtained as a byproduct from the DAST fluorination and allowed for the preparation of two biologically active compounds from a common precursor. Compound **1** and 2'-C-methylcytidine were assayed in a subgenomic HCV replicon assay system and found to be potent and selective inhibitors of HCV replication. Compound **1** shows increased inhibitory activity in the HCV replicon assay compared to 2'-C-methylcytidine and low cellular toxicity.

Introduction

Hepatitis C virus infection is a major health problem that leads to chronic liver disease, such as cirrhosis and hepatocellular carcinoma, in a substantial number of infected individuals. Once infected, about 20% of people clear the virus, but the rest can harbor HCV the rest of their lives. Ten to 20% of chronically infected individuals eventually develop liver-destroying cirrhosis or cancer. The current standard of care for chronic hepatitis C is combination therapy with an interferon-α and ribavirin. Studies have shown that more patients with hepatitis C respond to pegylated interferon-α/ribavirin combination therapy than to combination therapy with unpegylated interferon-α. The overall response rate to treatment, defined as loss of HCV from serum 6 months after completion of treatment, is 40%. Because of the low response rates as well as toxic side effects and unsustainable viral load reductions, these therapies are inadequate. Moreover, there is no established vaccine for HCV, and there is an urgent need for improved therapeutic agents that effectively combat chronic HCV infection.¹

The nonstructural protein NS5B has been characterized as an RNA-dependent RNA polymerase (RdRp) that is required for viral replication. This polymerase is considered to be an essential component in the HCV

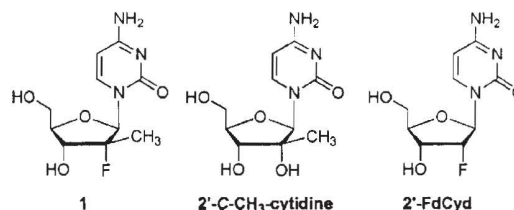


Figure 1. Structures of 2'-deoxy-2'-fluoro-2'-C-methylcytidine (**1**), 2'-C-methylcytidine, and 2'-deoxy-2'-fluorocytidine (2'-FdCyd).

replication complex and therefore is an ideal target for drug discovery. Recently, several 2'-modified nucleoside analogues with potent inhibitory activity against the HCV NS5B polymerase have been identified. Among the most potent compounds in this class are 2'-deoxy-2'-fluorocytidine (2'-FdCyd)² and 2'-C-methyl nucleosides (Figure 1).^{3–5} Despite the potent HCV inhibition of 2'-FdCyd, its therapeutic potential as an antiviral agent is diminished due to a lack of selectivity between host cells and the viral target. 2'-FdCyd triphosphate has been demonstrated to be a substrate for both RNA and DNA polymerases.^{6,7} Here we describe the synthesis and biological activity of 2'-deoxy-2'-fluoro-2'-C-methyl cytidine (**1**) as a potent anti-HCV agent.

Chemistry

For the synthesis of 2'-deoxy-2'-fluoro-2'-C-methylcytidine (**1**), *N*⁴-benzoyl-1-(2-methyl-3,5-di-*O*-benzoyl-β-D-arabinofuranosyl)cytosine (**6**) was chosen as the key intermediate and was prepared in approximately 20% yield in six steps from cytidine (Scheme 1).⁸ Briefly, selective benzylation of cytidine with benzoic anhydride in DMF,⁹ followed by treatment with TIDPSCl₂ in pyridine, afforded *N*⁴-benzoyl-3',5'-*O*-(tetraisopropyl-disiloxane-1,3-diyl)cytidine (**2**).¹⁰ Oxidation of the 2'-alcohol to the 2'-ketone derivative (**3**) was achieved with

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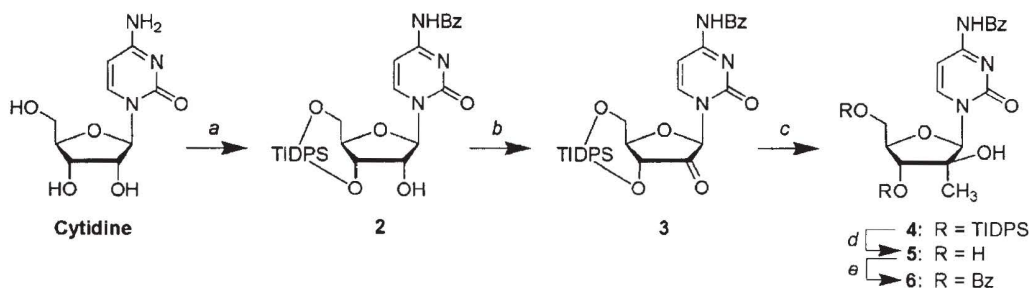
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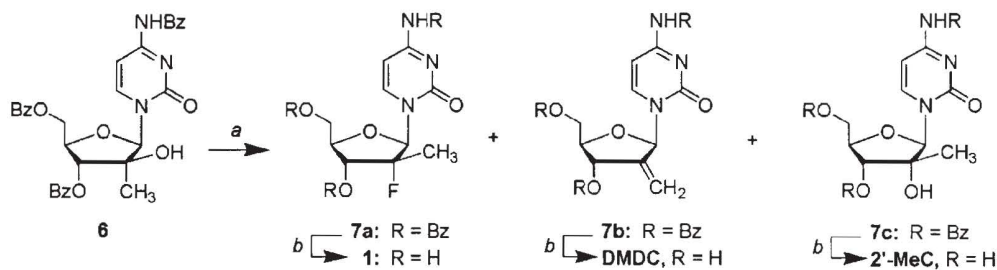
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Scheme 1^a

^a Reagents and conditions: (a) (i) Bz₂O, DMF, room temp, (ii) TIDPSCl₂, DMF; (b) DMSO, TFAA, TEA, -15 °C; (c) MeLi, -78 °C; (d) 1 M TBAF, concd HOAc, rt; (e) BzCl, pyridine, rt.

Scheme 2^a

^a Reagents and conditions: (a) DAST, toluene, -20 °C to rt; (b) MeOH/NH₃, rt.

Table 1. Fluorine-Coupled ¹H and ¹³C NMR Chemical Shifts, Multiplicities, and *J* Values for Compounds 1, 7a, 8, and 9^a

compd	2CH ₃	H-1/C-1	C-2	H-3/C-3
1	δ 1.17, d, ³ J _{H-F} = 22.3 δ 16.6, d, ² J _{C-F} = 25.9	δ 6.07, d, ³ J _{H-F} = 18.9 δ 88.6, d, ² J _{C-F} = 37.4	δ 101.2, d, ¹ J _{C-F} = 180.1	overlapping mult. δ 70.5, d, ² J _{C-F} = 18.3
7a	δ 1.49, d, ³ J _{H-F} = 22.4 δ 17.4, d, ² J _{C-F} = 25.2	δ 6.52, d, ³ J _{H-F} = 18.0 δ 91.2, d, ² J _{C-F} = 42.0	δ 100.2, d, ¹ J _{C-F} = 187.7	δ 5.56, dd, ³ J _{H-F} = 20.7 δ 72.4, d, ² J _{C-F} = 16.0
8	δ 1.39, d, ³ J _{H-F} = 22.3 δ 17.3, d, ² J _{C-F} = 25.1	δ 6.17, d, ³ J _{H-F} = 19.3 δ 90.7, d, ² J _{C-F} = 44.2	δ 99.9, d, ¹ J _{C-F} = 186.2	δ 5.49, d, ³ J _{H-F} = 21.2 δ 72.7, d, ² J _{C-F} = 16.1
9	δ 1.35, d, ³ J _{H-F} = 22.3 δ 16.9, d, ² J _{C-F} = 25.2	δ 6.13, d, ³ J _{H-F} = 18.9 δ 90.6, d, ² J _{C-F} = 44.0	δ 102.1, d, ¹ J _{C-F} = 180.1	overlapping mult. δ 72.5, d, ² J _{C-F} = 17.6

^a NMR spectra were recorded at 30 °C (400 MHz) in DMSO-*d*₆ for compound 1, CDCl₃ for compound 7a, CD₃OD + CDCl₃ for 8, and CD₃OD for 9 with concentrations of ~40 mg/0.75 mL. *J* values are in Hz.

trifluoroacetic anhydride/DMSO under Swern oxidation conditions.¹¹ Purification of compound 3 by silica gel chromatography followed by crystallization from petroleum ether-CH₂Cl₂ provided a white solid that was stable when stored at room temperature with minimal atmospheric exposure.¹² Treatment of the 2'-ketone (3) with methyl lithium at -78 °C in diethyl ether gave exclusively the protected 1-[2-*C*-methyl-3,5-*O*-(tetraisopropylidisiloxane-1,3-diyl)-β-D-arabinofuranosyl]cytosine (4).¹³ The 3',5'-silyl protecting group was removed with TBAF/acetic acid and replaced with benzoyl protecting groups to provide compound 6.

The fluorination of tertiary alcohols using DAST has been reported, but the stereochemistry of such transformations is substrate-specific and often unpredictable. For instance, Yang et al. reported that the DAST fluorination of a tertiary alcohol in 2-bromomethyl-DL-*myo*-inositol proceeds with retention of configuration.¹⁴ Wachtmeister et al. obtained a 4-fluoro-1-cyclopentanol containing a tertiary fluorine in 25% yield using DAST as a fluorinating reagent, and this transformation proceeded with inversion of configuration.¹⁵ Furthermore, dehydrations or eliminations, rearrangements, and ring contractions are often pervading problems in the DAST fluorination of highly functionalized mol-

Upon treating 6 with DAST in toluene or dichloromethane, a clean mixture of three products (7a-7c) in 15-20% yield for each compound was obtained (Scheme 2). The desired transformation of 6 to *N*⁴-benzoyl-1-[2-fluoro-2-methyl-3,5-di-*O*-benzoyl]cytosine (7a) proceeded with inversion of configuration and was stereospecific; no diastereomeric *N*⁴-benzoyl-1-[2-fluoro-2-methyl-3,5-di-*O*-benzoyl-β-D-arabinofuranosyl]cytosine was detected in the crude reaction mixture. The presence of the tertiary fluorine at the 2' position in 7a was confirmed by the ¹H and ¹³C NMR multiplicities and coupling constants (Table 1), whereas the stereochemistry of the fluorination was determined by nuclear Overhauser enhancement ¹H NMR difference spectroscopy (Figure 2).

Analysis of the ¹H NMR spectrum of compound 7a revealed three distinct multiplicities due to H-F coupling: a doublet at δ 1.49 (2'-CH₃), a doublet of doublets at δ 5.56 (H-3'), and a doublet at δ 6.52 (H-1'). Irradiation of the H-3' resonance resulted in a relatively large enhancement of both the H-5' signal (4.8%) and the 2'-methyl signal (5.9%), while irradiation of the 2'-methyl signal resulted in an enhancement of both H-3' (3.4%) and, to a lesser extent, H-1' (1.7%). Deprotection of 7a

(300 g) and chromatographed as before to give an off-white solid (46.4 g, 61%). Crystallization from aqueous acetone afforded an analytical sample: mp 197–200 °C; $[\alpha]_D^{25} + 132.0^\circ$ (c 1, MeOH); $^1\text{H NMR}$ (DMSO- d_6) δ 1.20 (s, 3H), 3.62–3.69 (m, 2H), 3.73–3.78 (m, 2H), 5.19 (t, 1H, $J = 5.4$ Hz), 5.25 (s, 1H), 5.52 (d, 1H, $J = 5.0$ Hz), 5.99 (s, 1H), 7.32 (d, 1H, $J = 5.8$ Hz), 7.50 (Ψ t, 2H, $J = 7.7$ Hz), 7.62 (Ψ t, 1H, $J = 7.3$ Hz), 8.00 (d, 2H, $J = 7.3$ Hz), 8.14 (d, 1H, $J = 6.9$ Hz), 11.22 (s, 1H); $^{13}\text{C NMR}$ (DMSO- d_6) δ 19.6, 61.3, 77.5, 78.5, 85.2, 88.9, 95.2, 128.5, 132.8, 133.3, 147.3, 154.9, 162.9, 167.4. Anal. Calcd (C₁₇H₁₉N₃O₆·0.5H₂O): C, 55.14; H, 5.41; N, 11.35. Found: C, 55.21; H, 5.47; N, 11.33.

N⁴-Benzoyl-1-(2-C-methyl-3,5-di-O-benzoyl- β -D-arabino-furanosyl)cytosine (6). Compound **5** (46.0 g, 0.127 mol) was dissolved in anhydrous pyridine (200 mL) and the solvent was removed in vacuo. The resulting syrup was dissolved in anhydrous pyridine, cooled to 0 °C under argon with stirring, and treated with BzCl (30.0 mL, 0.250 mol) dropwise over 10 min. After the addition was complete, the ice bath was removed and stirring was continued for 1.5 h. Water (5 mL) was added and the mixture was concentrated to dryness in vacuo. The residue was dissolved in CH₂Cl₂ and washed with saturated NaHCO₃ (1 × 500 mL) and water (1 × 500 mL). The organic layer was dried (Na₂SO₄), filtered, and concentrated to dryness to afford a syrup that was purified by silica gel chromatography eluting with 1:1 EtOAc–hexanes. Compound **6** was isolated as an off-white solid (48.5 g, 67%): $[\alpha]_D^{25} + 9.6^\circ$ (c 1, CHCl₃); $^1\text{H NMR}$ (CDCl₃) δ 1.64 (s, 3H), 4.50 (m, 1H), 4.78–4.85 (m, 2H), 5.50 (d, 1H, $J = 3.4$ Hz), 6.42 (s, 1H), 7.44–7.54 (m, 7H), 7.57–7.66 (m, 3H), 7.94 (d, 2H, $J = 7.8$ Hz), 8.05–8.09 (m, 4H), 8.21 (d, 1H, $J = 7.3$ Hz); $^{13}\text{C NMR}$ (CDCl₃) δ 19.9, 64.4, 79.2, 80.6, 81.6, 90.3, 96.7, 128.1, 128.6, 128.8, 128.9, 129.5, 129.9, 130.0, 133.2, 133.5, 134.0, 147.5, 156.0, 162.7, 165.9, 166.6. Anal. Calcd (C₃₁H₂₇N₃O₈): C, 65.37; H, 4.78; N, 7.38. Found: C, 65.59; H, 4.79; N, 7.16.

Reaction of 6 with DAST. To a stirred solution of **6** (2.96 g, 5.20 mmol) in anhydrous toluene (50 mL) was added DAST (1.0 mL, 7.8 mmol) at –20 °C under argon. After the addition was complete, the cooling bath was removed and stirring was continued for 1 h. The reaction mixture was poured into saturated NaHCO₃ (50 mL) and washed until gas evolution ceased. The organic phase was dried (Na₂SO₄), concentrated to dryness, and purified by silica gel chromatography eluting with 1:1:1 EtOAc–CHCl₃–hexanes to afford compound **7a** (0.55 g, 19%) as a white solid, followed by compound **7b** (0.39 g, 14%) as an off-white solid. Elution was continued with 1:1:1 EtOH–EtOAc–CHCl₃–hexanes to afford compound **7c** (0.451 g, 15%) as an off-white solid. Analytical samples were obtained by recrystallization from the indicated solvents.

N⁴-Benzoyl-3',5'-di-O-benzoyl-2'-fluoro-2'-C-methylcytidine (7a): mp 241 °C (CH₂Cl₂–hexanes); $[\alpha]_D^{25} + 82.0^\circ$ (c 1, CHCl₃); $^1\text{H NMR}$ (CDCl₃) δ 1.49 (d, 3H, $J = 22.4$ Hz), 4.64 (dd, 1H, $J = 3.44$, 12.9 Hz), 4.73 (d, 1H, $J = 9.5$ Hz), 4.90 (dd, 1H, $J = 2.4$, 12.7 Hz), 5.56 (dd, 1H, $J = 8.6$, 20.7 Hz), 6.52 (d, 1H, $J = 18.0$ Hz), 7.47–7.57 (m, 7H), 7.62–7.71 (m, 3H), 7.89 (d, 2H, $J = 6.9$ Hz), 8.07–8.11 (m, 5H), 8.67 (bs, 1H); $^{13}\text{C NMR}$ (CDCl₃) δ 17.4 (d, $J = 25.2$ Hz), 62.1, 72.4 (d, $J = 16.0$ Hz), 77.7, 91.2 (d, $J = 42.0$ Hz), 97.5, 100.2 (d, $J = 187.7$ Hz), 127.8, 128.6, 128.8, 128.9, 129.2, 129.6, 129.7, 130.3, 133.2, 133.4, 133.8, 134.1, 143.8, 154.6, 162.6, 165.6, 166.1; $^{19}\text{F NMR}$ (CDCl₃) δ 3.9 (m). Anal. Calcd (C₃₁H₂₆FN₃O₇·0.7H₂O): C, 63.74; H, 4.73; N, 7.19. Found: C, 63.71; H, 4.54; N, 7.20.

N⁴-Benzoyl-1-(2-deoxy-2-methylidene-3,5-di-O-benzoyl- β -D-glycero-pentofuranosyl)cytosine (7b): mp 173.4–174.4 (EtOH); $[\alpha]_D^{25} - 40.4^\circ$ (c 1, CHCl₃); $^1\text{H NMR}$ (CDCl₃) δ 4.58 (dd, 1H, $J = 3.7$, 5.0 Hz), 4.70–4.81 (m, 2H), 5.55 (s, 1H), 6.09–6.11 (m, 1H), 7.03 (d, 1H, $J = 1.3$ Hz), 7.40–7.66 (m, 10H), 7.85 (d, 1H, $J = 7.3$ Hz), 7.91 (d, 2H, $J = 7.7$ Hz), 8.03 (dd, 2H, $J = 0.9$, 8.3 Hz), 8.03 (dd, 2H), 8.86 (bs, 1H); $^{13}\text{C NMR}$ (CDCl₃) δ 63.9, 73.3, 80.7, 85.9, 97.8, 117.3, 127.8, 128.0, 128.7, 128.9, 129.1, 129.4, 129.6, 129.9, 130.2, 133.0, 133.6, 133.8, 144.1, 144.9, 155.1, 162.5, 165.9, 166.1. Anal. Calcd (C₃₁H₂₅N₃O₇): C, 67.51; H, 4.57; N, 7.62. Found: C, 67.21; H, 4.51;

N⁴-Benzoyl-2'-C-methyl-3',5'-di-O-benzoylcytidine (7c): mp 176.7–179.1 °C (EtOH); $[\alpha]_D^{25} + 46.2^\circ$ (c 1, CHCl₃); $^1\text{H NMR}$ (CDCl₃) δ 1.29 (s, 3H), 4.67–4.80 (m, 3H), 4.82–4.87 (m, 1H), 5.30 (d, 1H, $J = 5.8$ Hz), 6.09 (s, 1H), 7.47–7.56 (m, 4H), 7.89 (d, 2H, $J = 7.3$ Hz), 8.07–8.14 (m, 4H), 8.68 (s, 1H); $^{13}\text{C NMR}$ (CDCl₃) δ 21.4, 62.7, 75.8, 78.5, 79.0, 93.4, 97.3, 127.8, 128.5, 128.7, 128.9, 129.0, 129.4, 129.6, 130.0, 132.9, 133.2, 133.6, 144.3, 156.0, 162.8, 165.8, 166.2, 166.8. Anal. Calcd (C₃₁H₂₇N₃O₈·0.4H₂O): C, 64.56; H, 4.86; N, 7.29. Found: C, 64.54; H, 4.81; N, 7.32.

3',5'-Di-O-benzoyl-2'-deoxy-2'-fluoro-2'-methyluridine (8). Compound **7a** (0.225 g, 0.394 mmol) was suspended in 80% aqueous HOAc (15 mL) and heated under reflux with stirring for 12 h. The clear solution was cooled, concentrated to dryness in vacuo, and coevaporated with 50% MeOH–water (3 × 5 mL) to remove the residual HOAc. Purification by silica gel chromatography, eluting with 2% EtOH–CH₂Cl₂, gave 0.160 g of **8** (87%) as a white solid. Crystallization from 2-propanol afforded an analytical sample: mp 256.4–257.6 °C; $[\alpha]_D^{25} + 71.7^\circ$ (c 1, CHCl₃); $^1\text{H NMR}$ (CDCl₃ + CD₃OD) δ 1.39 (d, 3H, $J = 22.3$ Hz), 4.49 (dd, 1H, $J = 3.9$, 12.7 Hz), 4.57 (m, 1H), 4.79 (dd, 1H, $J = 2.7$, 12.5 Hz), 5.42 (d, 1H, $J = 8.1$ Hz), 5.49 (dd, 1H, $J = 9.20$, 21.2 Hz), 6.17 (d, 1H, $J = 19.3$), 7.37–7.50 (m, 4H), 7.51–7.57 (m, 3H, H-6), 7.93–8.01 (m, 4H); $^{13}\text{C NMR}$ (CDCl₃ + CD₃OD) δ 17.3 (d, $J = 25.1$ Hz), 62.1, 72.7 (d, $J = 16.1$ Hz), 90.7 (d, $J = 44.2$ Hz), 99.9 (d, $J = 186.2$ Hz), 103.1, 128.5, 128.6, 128.7, 129.4, 129.5, 130.1, 133.6, 134.0, 139.3, 150.4, 163.2, 165.7, 166.1; $^{19}\text{F NMR}$ (CDCl₃ + CD₃OD) δ 6.02 (m). Anal. Calcd (C₂₄H₂₁FN₂O₇): C, 61.54; H, 4.52; N, 5.98. Found: C, 61.42; H, 4.51; N, 5.96.

General Procedure for Deprotection. The free nucleosides were prepared by treating compounds **7a–c** and **8** with NH₃/MeOH (ca. 7 N, ~12 mL/mmol) followed by stirring at room temperature overnight (8–12 h). The solvent was removed in vacuo, and the compounds were isolated as indicated.

2'-Deoxy-2'-fluoro-2'-C-methylcytidine (1). Compound **7a** (6.30 g, 0.011 mol) was deprotected to give **1** (2.18 g, 76%) as a white powder after column chromatography eluting with 9% EtOH in CHCl₃ and then 17% EtOH and finally 25% EtOH in CHCl₃: mp 216.4–218.0 °C (EtOH); $[\alpha]_D^{25} + 125.6^\circ$ (c 1, H₂O); $^1\text{H NMR}$ (DMSO- d_6) δ 1.17 (d, 3H, $J = 22.3$ Hz), 3.63 (dd, 1H, $J = 2.7$, 13.7 Hz), 3.70–3.84 (m, 3H), 5.24 (app s, 1H), 5.60 (d, 1H, $J = 5.4$ Hz), 5.74 (d, 1H, $J = 7.71$ Hz), 6.07 (d, 1H, $J = 18.9$ Hz), 7.31 (s, 1H, NH₂), 7.42 (s, 1H, NH₂), 7.90 (d, 1H, $J = 7.3$ Hz); $^{13}\text{C NMR}$ (DMSO- d_6) δ 16.6 (d, $J = 25.9$ Hz), 58.5, 70.5 (d, $J = 18.3$ Hz), 81.4, 88.6 (d, $J = 37.4$ Hz), 94.4, 101.2 (d, $J = 180.1$ Hz), 140.5, 154.8, 165.2; $^{19}\text{F NMR}$ (DMSO- d_6) δ 2.60 (m). Anal. Calcd (C₁₀H₁₄FN₃O₄·1.5H₂O): C, 41.96; H, 5.94; N, 14.69. Found: C, 42.24; H, 5.63; N, 14.54.

Compound **1** was converted to the HCl salt and crystallized from aqueous ethanol: mp 243 °C (dec); $[\alpha]_D^{25} + 108.4^\circ$ (c 1, H₂O); $^1\text{H NMR}$ (DMSO- d_6) δ 1.29 (d, 3H, $J = 22.6$ Hz), 3.65 (dd, 1H, $J = 2.3$, 12.7 Hz), 3.76–3.90 (m, 3H), 5.96 (d, 1H, $J = 17.3$ Hz), 6.15 (d, 1H, $J = 7.9$ Hz), 8.33 (d, 1H, $J = 7.9$ Hz), 8.69 (s, 1.5H), 9.78 (s, 1.5H); $^{13}\text{C NMR}$ (DMSO- d_6) δ 16.2 (d, $J = 24.4$ Hz), 58.2, 69.9 (d, $J = 16.8$ Hz), 82.1, 88.8 (d, $J = 32.0$ Hz), 94.6, 101.1 (d, $J = 181.5$ Hz), 143.2, 147.6, 159.6; $^{19}\text{F NMR}$ (DMSO- d_6) δ 1.69 (m). Anal. Calcd (C₁₀H₁₅ClFN₃O₄): C, 40.62; H, 5.11; N, 14.21. Found: C, 40.80; H, 5.09; N, 14.23.

2'-Deoxy-2'-fluoro-2'-C-methyluridine (9). Deprotection of **8** (0.120 g, 0.209 mmol) followed by column chromatography eluting with 5–10% acetone in diethyl ether provided **9** (0.054 g, 100%) as a white solid: mp 237.3–238.0 °C; $[\alpha]_D^{25} + 83.2^\circ$ (c 1, MeOH); $^1\text{H NMR}$ (CD₃OD) δ 1.35 (d, 3H, $J = 22.3$ Hz), 3.79 (dd, 1H, $J = 2.1$, 12.5 Hz), 3.94–4.02 (m, 3H), 5.70 (d, 1H, $J = 8.1$ Hz), 6.13 (d, 1H, $J = 18.9$ Hz), 8.09 (d, 1H); $^{13}\text{C NMR}$ (CD₃OD) δ 16.9 (d, $J = 25.2$ Hz), 60.1, 72.5 (d, $J = 17.6$ Hz), 83.5, 90.6 (d, $J = 44.0$ Hz), 102.1 (d, $J = 180.1$ Hz), 103.0, 142.0, 152.4, 166.0; $^{19}\text{F NMR}$ (CD₃OD) δ 4.07 (bs). Anal. Calcd (C₁₀H₁₃FN₂O₅): C, 46.16; H, 5.04; N, 10.77. Found: C, 45.96; H, 4.93; N, 10.49.

Isolation of 2'-C-methylcytidine. Compound **7c** (0.1 g,

gave a white solid (0.032 g, 71%): mp 244.2–245.8 °C (lit.¹³ mp 239.5–242 °C, lit.¹⁸ mp 243–245 °C); $[\alpha]_D^{23} +135.7^\circ$ (c 1, H₂O) (lit.¹⁸ $[\alpha]_D +132^\circ$ (c 0.5, H₂O), lit.¹⁹ $[\alpha]_D^{20} +128^\circ$ (c 1, H₂O)); ¹H NMR (DMSO-*d*₆) δ 0.92 (s, 1H), 3.58–3.62 (m, 2H), 3.70–3.77 (m, 2H), 4.98 (s, 1H), 5.06 (d, 1H, *J* = 7.3 Hz), 5.11 (t, 1H, *J* = 5.0 Hz), 5.67 (d, 1H, *J* = 7.7 Hz), 5.87 (s, 1H), 7.11 (s, 1H), 7.17 (s, 1H), 7.94 (d, 1H, *J* = 7.2 Hz); ¹³C NMR (DMSO-*d*₆) δ 20.0, 59.0, 71.8, 78.3, 81.8, 91.3, 93.6, 141.1, 155.5, 165.4.

Isolation of 2'-Deoxy-2'-methylidene cytidine (DMDC). Compound **7b** was deprotected and crystallized from H₂O: mp 190–194 °C (dec) (transition at 90–92 °C) (lit.²⁰ mp 89–90 °C); $[\alpha]_D^{22} -39.8^\circ$ (c 1, H₂O); ¹H NMR (DMSO-*d*₆) δ 3.50–3.70 (m, 3H), 4.43 (broad t, 1H), 4.93 (t, 1H, *J* = 5.2 Hz), 5.13 (s, 1H), 5.29 (s, 1H), 5.61 (d, 1H, *J* = 6.4 Hz), 5.69 (d, 1H, *J* = 7.6 Hz), 6.51 (s, 1H), 7.20 (s, 1H), 7.23 (s, 1H), 7.47 (d, 1H, *J* = 7.6 Hz); ¹³C NMR (DMSO-*d*₆) δ 60.5, 69.8, 84.1, 84.2, 94.6, 110.8, 142.0, 151.1, 155.3, 165.5.

Biological Methods. Antiviral assays with bovine viral diarrhea virus and the HCV replicon were performed as described previously.¹⁷

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