Stability of Saturated and Unsaturated Perfluoroalkyl Telomer Acid Compounds as Reference Standards Gilles Arsenaulta, Brock Chittima, Alan McAleesa, Robert McCrindleb, Nicole Riddella, Gregg Tomvo and Brian Yeoa, ^a Wellington Laboratories Inc., Research Division, Guelph, Ontario, Ontario, N1G 3M5, Canada ^b University of Guelph, Chemistry Dept., Guelph, Ontario, N1G 2W1, Canada ^c Dept. Fisheries and Oceans, Winnipeg, Manitoba, R3T 2N6, Canada Introduction Results and Discussions (continued) **Results and Discussions (continued)** · Fluorotelomer alcohols are a group of compounds used in various commercial Interestingly, the type of glass in which the solutions are stored was also shown to products from which they may escape into the environment and then get oxidized or Since the rate of dehydrofluorination of the saturated telomer acids in the presence have an effect on the stability of the saturated telomer acids (see figure 5). metabolized to fluorotelomer acids. of HF was still unacceptable, more studies were conducted to find conditions to These results indicate that the basicity of the glass has an impact on the To fully understand what is actually occurring in the environment, accurate analysis of further reduce the rate of this decomposition. dehydrofluorination rates of the saturated telomer acids. fluorotelomer alcohols and acids in environmental samples is needed and this A study comparing the effects of different acids in methanol showed that requires the use of reference standard solutions.¹ Figure 5. Effect of glass type on Figure 6. Rate of dehydrofluorination The facile degradation of saturated to unsaturated telomer acids has been reported.² trifluoroacetic acid (TFA) and HCl were the best of those tested in stabilizing the in IPA with/without HCI and dehydrofluorination rates in MeOH saturated telomer acids (see figure 3). Objective at 22°C or 4°C at ambient temperature (22°C) • The objective of this work was to examine carefully the stability of these telomer acids Figure 3a. Comparison of Figure 3b. Comparison of Effect of plass typ in solutions when prepared for use as reference standards in analytical methods. dehydrofluorination rates for dehvdrofluorination rates for This work may provide further insight into the behaviour of these telomer acids in the FOEA in the presence of FOEA in the presence of environment. various acids at 22°C various acids at 4°C Experimental LC/MS: LC/MS experiments were conducted on a Waters Acquity Ultra Performance LC attached to a Micromass Quattro micro API. Separations were performed on an Acquity UPLC BEH Shield RP18 column (1.7 um, 2.1 x 100 mm). Typical run conditions were 65% (80:20 MeOH; ACN) and 35% water, both with 10 mM NH4OAc at a flow rate of 300 uL/minute. Time (davs The 8:2 saturated telomer acid (FOEA) was used in this study. Solutions prepared in IPA containing a small amount of HCI show very good stability Results and Discussions against dehydrofluorination (see figure 6). As expected, solutions without HCI The conversion of a saturated to an unsaturated telomer acid is easily achieved in the showed poor stability. Solutions without HCI and in clear borosilicate glass showed presence of a base such as sodium hydroxide³ (see figure 1). the highest rates of dehydrofluorination. Figure 1. Dehydrofluorination of a saturated telomer acid (FOEA) to an unsaturated telomer acid (FOUEA) Figure 4. Comparison of Comment on the stability of unsaturated telomer acids C₆F₁₃-CF₂-CF₂-CH₂-CO₂H HF dehydrofluorination It is known⁵ that unsaturated telomer acids can react with MeOH in the presence of a rates in various base to form methoxy-substituted unsaturated telomer acids (see Figure 7). solvents at ambient FOUEA FOFA Figure 7. Reaction of FOUEA with methanol temperature (22°C) Initial studies in methanol indicated that the presence of HF had a stabilizing effect for CeF12-CF2 FOEA but the elimination process was still occurring (see figure 2). MeOH + HF Figure 2. Comparison of The stabilization trend observed of HCI ~ TFA > acetic acid > HF indicates that FOUEA MeO-FOUEA dehydrofluorination decreased basicity of the anion leads to a decrease in dehydrofluorination. No HF & 20°C rates for FOEA with A comparison of different solvents showed that isopropanol (IPA) was the best LC varying amounts of Indeed, when FOUEA is stored in methanol at a concentration of 50ug/ml, formation solvent for stabilizing the saturated telomer acid (see figure 4). HE in Methanol of the methoxy adduct does occur as evidenced by LC/MS. This formation is slow, with conversion being measured at about 1% per year at ambient temperature. y = 1238-064 Storage at 4°C is recommended for solutions of these unsaturated telomer acids in It appears that the dielectric constant of the solvent may be a factor that influences methanol in order to further reduce the rate of formation of the methoxy-adduct. the stability of the saturated telomer acid (see Table 1). It is known that fluoride anion can behave as a weak base⁴. Thus, the addition of HF, while rendering the medium acidic, also introduces fluoride anion which could itself Table 1. Dielectric constants of the various solvents act as a dehydrofluorination agent. Dielectric Constant Solvent Type Acetonitrile (ACN) Aprofic 36.6 Conclusions References Methanol (MeOH) Protic 33 ¹Martin JW, Kannan K, Berger U, De Voogt P, Field J, Franklin J, Giesy JP, Harner T, Muir DCG, Solutions of saturated telomer acids are best kept in IPA in the presence of trace Scott B, Kaiser M, Jarnberg U, Jones KC, Mabury SA, Schroeder H, Simcik M, Sottani C, Isopropanol (IPA) Protic 18.3 amounts of HCI Van Bavel B. Karrman A. Lindstrom G and Van Leeuwen S. Environ. Sci. Technol. 2004: 38. Tetrahydrofuran (THF) Aprotic 7.5 2484-2554 Analytical laboratories have to be careful how they handle both reference standard Methyl tert-butyl ether (MTBE) Aprotic 4.5 ²Loewen M, Halldorson T, Wang F and Tomy G. Environ. Sci. Technol. 2005; 39, 2944-2951. solutions and environmental samples if they wish to analyze for saturated telomer Toluene Aprotic 2 ³Achilefu S, Mansuy L, Selve C and Thiebaut. J.Fluorine Chem. 1995; 70, 19-26.

⁴Clark JH, Chem, Rev. 1980; 80, 429-452.

⁵Zuczek C, Gerardin-Charbonnier C, Rocca S, Thiebaut S and Selve C, J. Fluorine Chem, 1999; 99, 41-49,

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