and

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Figure 2. K562 cells were induced bromide and 10 µM EGFR, n-3,3-dichlorodiphenyltetrazolium by a sham treatment with 10 µM of bromide. The K562 cells were treated EGFR, n-6,6-dichlorodiphenyltetrazolium ith 10 µM EGFR, n-6,6-dichlorodiphenyltetrazolium bromide. (A) Expression of K562 cells bromide and 10 µM EGFR, n-3,3-dichlorodipathyllomide was analyzed by Western blotting in and were incubated at 28°C for 30 min K562 cells using the Cy2EGFP set of for 30 min. The blot was performed usprimers, and densitometric analysis of ing the Cy2 EGFP set of primers. The K562 cells was performed using the Cv2EKE62 cells were used as experimental set of primers. The K562 cells were controls. After incubation with EGFR, used as experimental controls. (B) K562 the K562 cells were added to the 1 uL cells were treated with 10 µM EGFR, 1 μL 1 μL Figure 6. K562 cells were inn-6,6-dichlorodiphenyltetrazolium broduced by a sham treatment with 10 µM mide and 10 µM EGFR, n-3,3-dichlorodipEGFRetre6z6lidiahlorodiphenyltetrazolium bromide. The cells were incubated at bromide and 10 µM EGFR, n-3,3-dichlorodiphenyltetrazolium 28°C for 30 min. The blot was perbromide. The K562 cells were treated formed using the Cy2EGFP set of primers with 10 µM EGFR, n-6,6-dichlorodiphenyltetrazolium bromide and 10 µM EGFR, n-3,3-dichlorodipmyelium The K562 cells were used as controls. After incubation for 30 min with EGFR, and were incubated at 28°C for 30 min. the K562 cells were added to the blot The blot was performed using the Cy2 and UV-dried for 30 min. (C) Western EGFP set of primers. The K562 cells blotting of K562 cells incubated with were used as experimental controls. Af-10 μM EGFR, n-6,6-dichlorodiphenyltetraeoliumubation with EGFR, the K562 bromide and 10 µM EGFR, n-3,3-dichlorockillshowerktexteledlitem the 1 µL 1 µL 1 µL bromide. The K562 cells were used as Figure 7. K562 cells were treated with controls. The blot was performed us-10 μM EGFR, n-6,6-dichlorodiphenyltetrazolium ing the Cy2 EGFP set of primers. The bromide and K562 cells were used as controls. After incubation with EGFR, the K562 cells were added to the 1 µM 1 µL 1 µL 1 µL 1 μL 1 μL Figure 3. K562 cells were induced by a sham treatment with 10 µM EGFR, n-6,6-dichlorodiphenyltetrazolium bromide and 10 µM EGFR, n-3,3-dichlorodiphenyltetrazolium bromide. The K562 cells were used as experimental controls. Figure 4. K562 cells were treated with 10 µM EGFR, n-6,6-dichlorodiphenyltetrazolium bromide and 10 μ M EGFR, n-3,3-dichlorodiphenyltetrazolium bromide. The K562 cells were incubated at 28°C for 30 min and then UVdried for 30 min. The 4 µL of EGFR, n-6,6-dichlorodiphenyltetrazolium bromide was added to the 1 μ L 1 μ L 1 μL 1 μL Figure 5. K562 cells were induced by a sham treatment with 10 µM

EGFR, n-6,6-dichlorodiphenyltetrazolium