# Publications Using GAIN Genotyping Data

1. Aulchenko, Y.S., Ripatti, S., Lindqvist, I., Boomsma, D., Heid, I.M., Pramstaller, P.P., Penninx, B.W., Janssens, A.C., Wilson, J.F., Spector, T., Martin, N.G., Pedersen, N.L., Kyvik, K.O., Kaprio, J., Hofman, A., Freimer, N.B., Jarvelin, M.R., Gyllensten, U., Campbell, H., Rudan, I., Johansson, A., Marroni, F., Hayward, C., Vitart, V., Jonasson, I., Pattaro, C., Wright, A., Hastie, N., Pichler, I., Hicks, A.A., Falchi, M., Willemsen, G., Hottenga, J.J., de Geus, E.J., Montgomery, G.W., Whitfield, J., Magnusson, P., Saharinen, J., Perola, M., Silander, K., Isaacs, A., Sijbrands, E.J., Uitterlinden, A.G., Witteman, J.C., Oostra, B.A., Elliott, P., Ruokonen, A., Sabatti, C., Gieger, C., Meitinger, T., Kronenberg, F., Doring, A., Wichmann, H.E., Smit, J.H., McCarthy, M.I., van Duijn, C.M. and Peltonen, L. (2009) Loci influencing lipid levels and coronary heart disease risk in 16 European population cohorts, Nat Genet, 41, 47-55.

2. Estrada, K., Krawczak, M., Schreiber, S., van Duijn, K., Stolk, L., van Meurs, J.B., Liu, F., Penninx, B.W., Smit, J.H., Vogelzangs, N., Hottenga, J.J., Willemsen, G., de Geus, E.J., Lorentzon, M., von Eller-Eberstein, H., Lips, P., Schoor, N., Pop, V., de Keijzer, J., Hofman, A., Aulchenko, Y.S., Oostra, B.A., Ohlsson, C., Boomsma, D.I., Uitterlinden, A.G., van Duijn, C.M., Rivadeneira, F. and Kayser, M. (2009) A genome-wide association study of northwestern Europeans involves the CNP signaling pathway in the etiology of human height variation, Hum Mol Genet. Available on-line only.

3. Pardo, L., Bochdanovits, Z., de Geus, E., Hottenga, J.J., Sullivan, P., Posthuma, D., Penninx, B.W., Boomsma, D. and Heutink, P. (2009) Global similarity with local differences in linkage disequilibrium between the Dutch and HapMap-CEU populations, Eur J Hum Genet, 17, 802-810.

4. Shi, J., Levinson, D.F., Duan, J., Sanders, A.R., Zheng, Y., Pe'er, I., Dudbridge, F., Holmans, P.A., Whittemore, A.S., Mowry, B.J., Olincy, A., Amin, F., Cloninger, C.R., Silverman, J.M., Buccola, N.G., Byerley, W.F., Black, D.W., Crowe, R.R., Oksenberg, J.R., Mirel, D.B., Kendler, K.S., Freedman, R. and Gejman, P.V. (2009) Common variants on chromosome 6p22.1 are associated with schizophrenia, Nature, 460, 753-757.