



# The high-resolution G-banded karyotype of *Sus scrofa domestica* L

O Galman, Martine M. Yerle, G Echard

## ► To cite this version:

O Galman, Martine M. Yerle, G Echard. The high-resolution G-banded karyotype of *Sus scrofa domestica* L. *Genetics Selection Evolution*, 1991, 23 (Suppl1), pp.113s-116s. hal-00893912

**HAL Id: hal-00893912**

**<https://hal.science/hal-00893912>**

Submitted on 11 May 2020

**HAL** is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

## The high-resolution G-banded karyotype of *Sus scrofa domestica* L

O Galman, M Yerle, G Echard

*Institut National de la Recherche Agronomique,  
Laboratoire de Génétique Cellulaire, BP 27, 31326 Castanet-Tolosan, France*

(Proceedings of the 9th European Colloquium on Cytogenetics of Domestic Animals;  
Toulouse-Auzeville, 10-13 July 1990)

**high resolution banding / karyotype / pig / GTG-banding**

The need for a high-resolution G-banded karyotype of the pig has been demonstrated following the publication by the Committee for a Standardized Karyotype of *Sus scrofa* (CSKSS, 1988) of its standard karyotype which was based on moderately extended G- and R-banded chromosomes. Rønne *et al* (1987) had presented a high-resolution R-banded karyotype at the 541 band level but no corresponding G-banded karyotype has been published to date. To fill this gap, this paper describes a high-resolution GTG-banded pig karyotype at the 539 band level.

To obtain mitotic spreads at late prophase, early and mid-metaphase stages, pig lymphocytes were synchronized with methotrexate ( $10^{-7}$  M) for 18 h to block cells at S phase, then subsequently released by leucovorin ( $3 \times 10^{-4}$  M) and thymidine ( $10^{-5}$  M). Ethidium bromide ( $2.5 \times 10^{-5}$  M) and colcemid ( $5 \times 10^{-7}$  M) were employed 2 and 0.5 h, respectively, before harvest. Hypotonic treatment, GTG-banding, photography and idiogram construction have been described elsewhere (Yerle *et al*, 1987).

The evolution of G-bands of each of the 38 pig chromosomes was analyzed from photographs of 52 well-spread and banded mitoses at progressive mitotic stages from metaphase (CSKSS standard) to late prophase. The final idiograms of the chromosomes with 4 haploid karyotypes at the 539 band level, shown in figure 1, take into consideration all 160 positive, 278 negative and 101 intermediate bands and subbands, according to their relative positions and staining intensities (Yerle *et al*, 1991). Using standard landmarks and nomenclature of major bands as reference points, the fate of each band was studied. For each chromosome, 3 or 4 intermediate stages were constructed to indicate which bands had subdivided, appeared or were retained in the final elongated stage. In most long chromosomes (*eg*, 1, 4, 6, 8, 9, 13, 14 and X) landmarks established to recognize the chromosomes are no longer evident, for example band q.4.1 on chromosome 13. However, in other chromosomes some bands, *eg*, q.2.1.1 of chromosome 1, are still distinct in the definitive stage (fig 1). The persistence of centromeric dark bands in the telocentric chromosomes

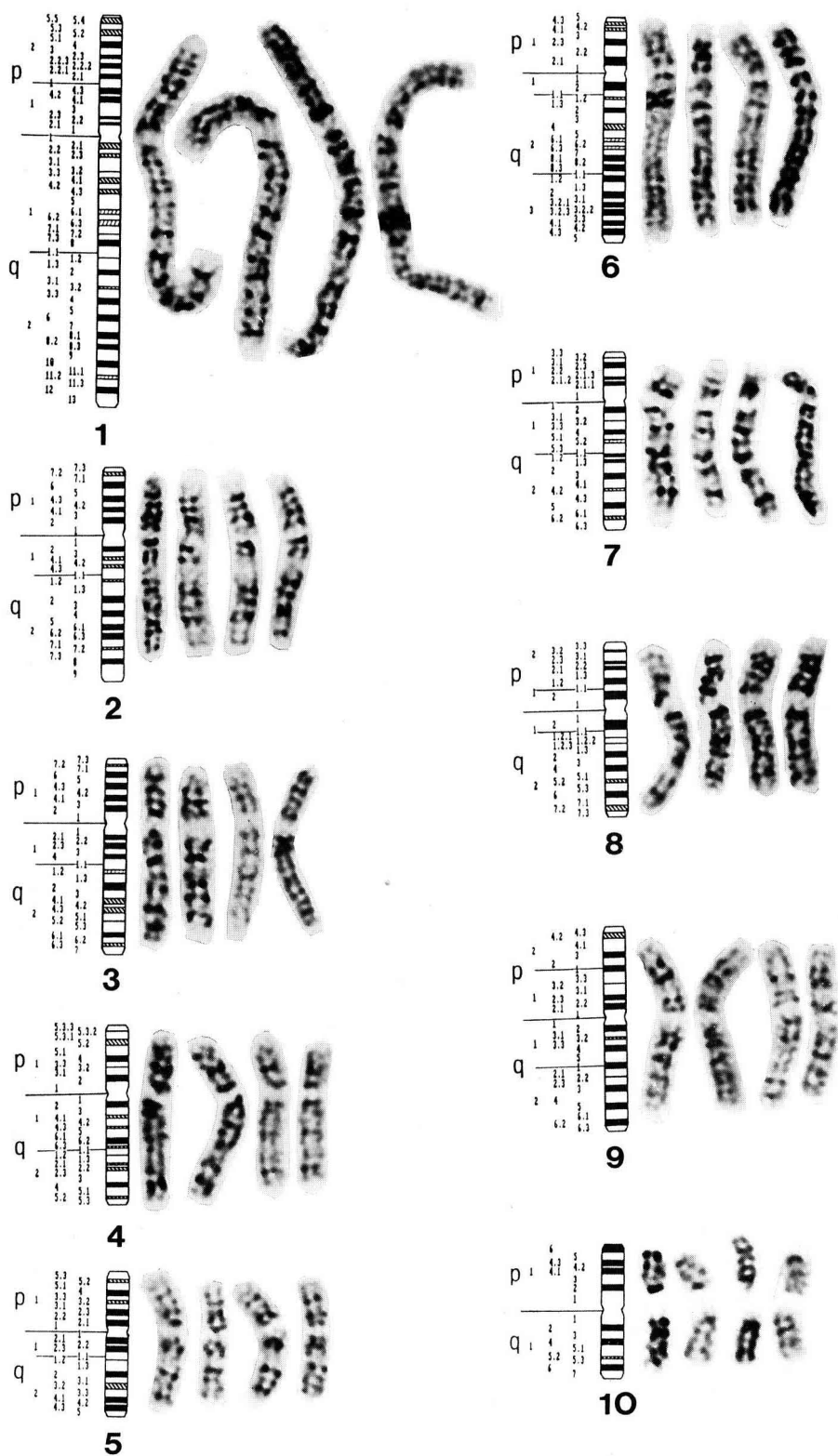


Fig 1. High-resolution GTG-banded chromosomes of the pig and their idiograms in the late prophase stage. The chromosomes represent composite sets from numerous cells. The idiograms, based on band measurements and on relative staining intensities, were made with the aid of an IBM PC computer. The longest horizontal lines indicate the position of the centromere.

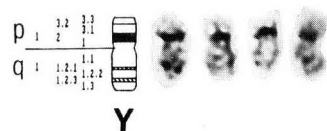
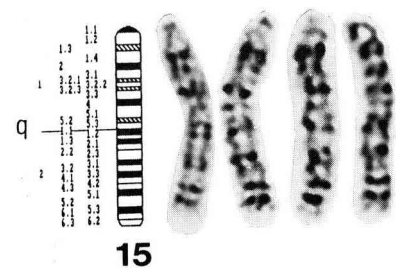
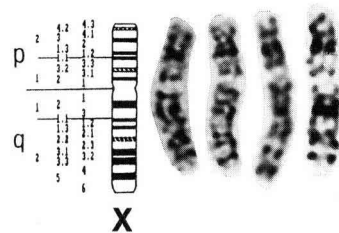
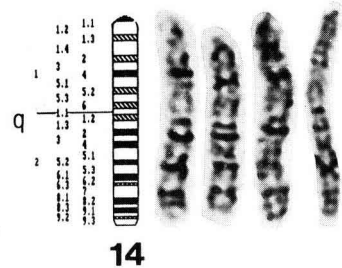
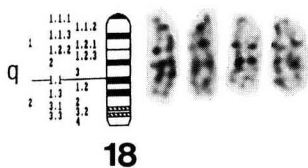
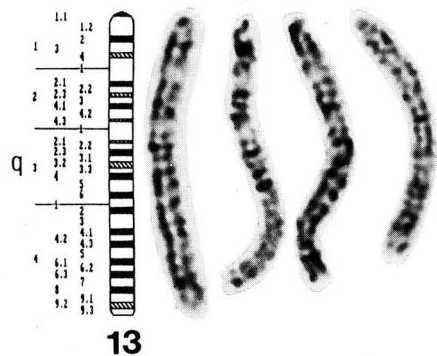
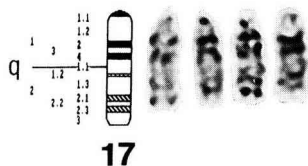
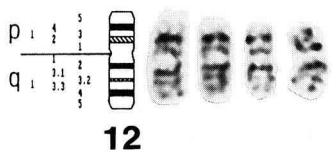
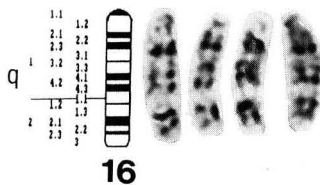
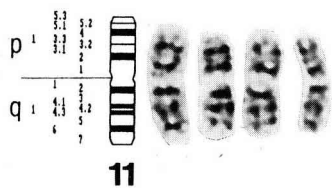


Fig 1. Continued

(13–18) from the standard to the final stage should also be noted. These bands, which correspond to (or overlap) the terminal R+ bands (Rønne *et al*, 1987), are likely to be the centromeric heterochromatic C-bands which were also stained during GTG-banding.

The karyotype proposed here can be a useful tool for the study of comparative chromosome organization and the precise mapping of the porcine genome (Yerle *et al*, 1990).

## REFERENCES

- Committee for the Standardized Karyotype of *Sus scrofa*. (1988) Standard karyotype of the domestic pig. *Hereditas* 109, 151-157
- Rønne M, Poulsen BS, Shibasaki Y, Flou S, Elberg JJ (1990) The high-resolution R-banded karyotype of the domestic pig *Sus scrofa domestica* L. *Cytobios* 49, 103-109
- Yerle M, Echard G, Gillois M (1987) The high resolution GTG-banding pattern of rabbit chromosomes. *Cytogenet Cell Genet* 45, 5-9
- Yerle M, Gellin J, Dalens M, Galman O (1990) Localization on pig chromosome 6 of markers: GPI, APOE, ENO1 carried by chromosomes 1 and 19, using *in situ* hybridization. *Cytogenet Cell Genet* 54, 86-91
- Yerle M, Galman O, Echard G (1991) The high resolution G-T-G banding pattern of pig chromosomes. *Cytogenet Cell Genet* 56, 45-47