Ammotragus is derived from the ancestral caprine form by one centric fusion. Furthermore, the identity between Ammotragus and Ovis vignei karyotypes supports GEIST's³ view that an Ammotragus-like form was directly ancestral to Ovis. Finally, the chromosomal homologies in the series of 1 to 3 biarmed autosomal pairs in Ovis lend further strength to the argument that Ovis vignei (2n = 58) is descended from the ancestral Ovis population, and that 2n = 54 Ovis are the most recently evolved of wild sheep 9 , 11 .

The fossil record indicates that *Ovis* and *Capra* have been separated for more than a million years 4,5 . During that time, their chromosomes exhibited structural stability, evolutionary change consisting merely of whole-arm rearrangements. Morphological 2 and behavioral 3 differences between sheep and goats are thus due to the accumulation of gene differences. The cytogenetic evidence suggests there are two main lineages among surviving caprines. Starting from a hypothetical rupicaprine-type ancestor with a primitive 2n = 60, FN = 60 karyotype, one lineage evolved through an intermediate,

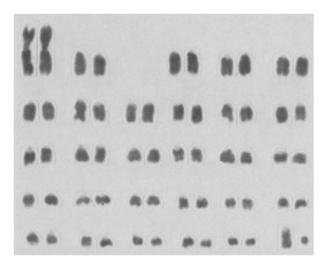


Fig. 4. Karyotype of a male Ovis vignei (2n = 58). The biarmed autosomes conform to the pattern of pair I.

aoudad-like form to the true sheep, with reductions in diploid number. In the true goat (Capra) lineage, morphological differentiation proceeded while the karyotype remained conservative; in contrast, the thar (Hemitragus) has remained morphologically close to the hypothetical rupicaprine ancestor3 while its chromosome number has been reduced $(2n = 48)^7$. The 5th member of the tribe Caprini, the bharal (Pseudois nayaur) has a reduced chromosome number $(2n = 54)^{13}$ but morphologically exhibits convergence toward true sheep2,3. G-bands in Hemitragus and Pseudois must be studied to confirm their chromosomal relationship to other caprines and it will be of interest to determine whether the biarmed chromosomes of Pseudois are homologous to those of 2n = 54 Ovis. If so, it would support the concept of convergent evolution at the chromosomal level.

ВЫВОДЫ. Хромосмы и Γ -полосы диких баранов, Африканских гривистых баранов, и безоаровых козлов изучалы. Сарта aegagrus из Ирана показало 2n=60, и кариотип содержит 29 пары акроцентических аутосом. Шесть из этих аутосомов одинаковый с трёх пар двеплечных аутосом у диких баранов и домашных овец (2n=54). Аmmotragus Iervia и Ovis vignei (2n=58) имели кариотипы и Γ -полосы пар две-плечных аутосом неразличимый.

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Effects of Abscisic Acid on Photosynthesis of Protoplasts from Petunia hybrida

Protoplasts from plants are suitable experimental objects for investigations in which substances are taken up by cells, since chemicals can reach the cell surface directly. Therefore, it is convenient to use them for experiments concerned with effects of phytohormones on metabolism. So far, some work on the application of auxins 1-3 has been described, and recently we have made some investigations on the influence of zeatin on photosynthesis and respiration of isolated mesophyll protoplasts from *Petunia* (Kull and Hoffmann, in preparation).

Application of abscisic acid (ABA) leads to a decrease of photosynthetic rate in treated leaves, very likely due to stomatal closure caused by the phytohormone 4-6. Some observations 7 suggest that mesophyll photosynthesis is not affected directly. Recently, it was found that in illuminated and unilluminated dark grown Avena seedlings and in etio-chloroplasts of the same species, ABA reduced the levels of ribulosediphosphate carboxylase

activity. We studied the effect of ABA on the photosynthesis of mesophyll protoplasts from *Petunia*. The influence of ABA on the respiration of these isolated protoplasts was briefly investigated in connection with our work on zeatin.

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Effect of abscisic acid on		

Time ABA acting (h)	Concentration of ABA (racemate)						
	1 ppm 3.8 μM/l	5 ppm 19μM/l	10 ppm 38 μM/l	50 ppm 190 μM/l	200 ppm 76 0 μM/l		
						1	100 + 4.5
2	100 + 4.5	100 ± 4.5	104 ± 4.5	131 ± 3.0	-		
3	100 ± 4.5	103 ± 4.5	123 ± 3.0	198 ± 3.0	116 ± 3.0		

Controls = 100%.

The mesophyll protoplasts from Petunia hybrida were produced by methods already described 9, 10. In scintillator vials 106 protoplasts in 1 ml medium (0.4 M mannitol/ 0.23 M NaHCO₃, ratio 95:5; with 0.22 M HCl brought to pH 7.2) were incubated with ABA (racemate) of different concentrations (Table) for different times (1-3 h). Then $2~\mu\text{Ci}$ of NaH $^{14}\text{CO}_3$ in 100 μl 0.4 M mannitol were added and the probes were illuminated for exactly 10 min in an illuminated thermostate (8×40 W Osram-L-Warmton lamps, ca. 9000 lux, temperature 24°C). After this, hot ethanol was added and the vials boiled for 2 min. After transferring into 50 ml graduated vessels and filling with 60% ethanol, the liquid was boiled again for 30 min, then the vessels filled up with ethanol. The activity of an equivalent of this final solution was measured in the scintillation counter in dioxane cocktail (gain 83%). Since ABA must be brought in solution with a few droplets of ammonia, the concentration of the latter rises with increasing concentration of the hormone. Therefore the respective concentrations of NH₃ were established in the controls. Always after addition of ABA or - in the controls - of NH₃ alone, the pH was adjusted again to 7.2. Due to the rising quantities of NH_a, the absolute rates of photosynthesis, measured as amount of ¹⁴CO₂ fixed, are not comparable; in the controls they decline. (In 1 set of control experiments without any NH₃ added, the rate is 2.98 μM CO₂/mg chlorophyll in 10 min; this is in good agreement with our earlier findings). Therefore the results are given as percentage of the photosynthetic rate of the respective controls. Each experiment was done 3 times together with 3 controls.

The results are shown in the Table. A rise of the photosynthetic rate caused by ABA can be seen. A marked effect needs relatively high concentrations of ABA with respect to naked protoplasts. With $1.9 \times 10^{-4}~M/l$ ABA after 3 h the increase of the photosynthesis reaches nearly 100%; with $7.6 \times 10^{-4}~M/l$ the effect is considerably lower, probably due to an inhibitory effect of very high concentrations of the phytohormone.

To a certain extent the results are surprising, considering that ABA is a senescence-promoting hormone ^{11, 12}. The concentrations of ABA which cause a rise of the photosynthetic rate of mesophyll protoplasts are a great deal higher than those needed for closure of stomata and the subsequent inhibition of photosynthesis of

leaves ⁶. Therefore an unspecific effect of ABA cannot be fully excluded. The results show the effects of a phytohormone on isolated cells in this case to be different from those on the whole leaf as a target. Under normal physiological conditions ABA is effective in whole tissues, so in evolution the function of the phytohormone was determined by selection in respect to the whole system and not to single cells. We think the results are explained best, in agreement with the findings of some other authors, as an effect of ABA on membrane permeability ^{4,13}, or the activity of ion uptake ^{4,14–16}. The reported ⁸ reduction of the activity of ribulosediphosphate carboxylase activity is due to an inhibition of plastid development and is not contrary to our findings with 'adult' cells ¹⁷.

Zusammenfassung. Inkubation mit Abscisinsäure verursacht bei Mesophyll-Protoplasten von Petunia hybrida einen Anstieg der Photosyntheserate.

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Der Einfluss der Aggression auf Lernen und Vergessen beim Kampffisch Betta splendens (Pisces, Anabantoidei)

Lern- und Behaltensprozesse werden stark von allem beeinflusst, was das lernende Individuum vor, während und nach dem Lernen tut und erlebt. Konsolidierung und Stabilität der Engramme sind offensichtlich am grössten, wenn auf den Lernakt Perioden absoluter Passivität, etwa Schlaf, oder aber Ruheperioden folgen ¹⁻⁹. Zwischen Lernen und Reproduzieren eingeschobene Aktivitäten scheinen dagegen den Vergessensvorgang zu beschleuni-