Disorders of Neuronal Migration

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ABSTRACT: Neuronal migration constitutes one of the major processes by which the central nervous system takes shape. Detailed knowledge about this important process now exists for different brain regions in rodent and monkey models as well as in the human. In the human, distinct genetic, chromosomal and environmental causes are known that affect neuronal migration, often in a morphologically distinct pattern, but the underlying pathological mechanisms are largely unknown. This review is intended to integrate our basic knowledge of the field with the accumulated intelligence on a large number of disorders and syndromes that represent the human part of the story.

RéSUMÉ: Perturbations de la migration neuronale La migration neuronale constitue un des processus les plus importants par lequel le système nerveux central est façonné. Nous possédons actuellement des connaissances détaillées sur ce processus important dans différentes régions du cerveau de modèles animaux (rongeurs et singes) ainsi que chez l’humain. Chez l’humain, des causes génétiques, chromosomiques et environnementales distinctes sont connues comme affectant la migration neuronale, donnant lieu à des patterns morphologiques souvent distincts; les mécanismes pathologiques sous-jacents sont pour la plupart inconnus. Dans la présente revue, nous désirons intégrer nos connaissances de base dans ce domaine avec les données accumulées au sujet d’un grand nombre d’affections et de syndromes représentant leur contrepartie chez l’humain.


After the closing of the neural tube and the formation of the telencephalic vesicles, neuronal migration is the main process by which topical differentiation within the brain is effected. By this process many billions of newly generated neural cells are addressed to their proper position mainly in nuclear masses or in the cerebral and cerebellar cortices. General or topical loss of control over this process is generally called neuronal migration disorder (abbreviated NMD). NMD will result in either cell death or improper positioning of functional cell groups. This in turn will result in failing connections or improper wiring (misconnections) responsible for functional deficiencies and epilepsy. The clinical relevance of NMD is highlighted by an increasing body of literature on a number of specific clinical entities, either inherited or prenatally acquired, and by the increasing resolution of imaging techniques by which NMD can be detected or at least suspected. Basic understanding of neuronal migration, mainly by morphological observations on rodent embryos, either normal or belonging to strains harboring inherited NMD, has increased substantially over the past twenty years; the understanding of the process in biochemical terms is emerging. NMD in the human embryo-fetus may arise from monogenetic (metabolic), chromosomal, hypoxic-ischemic and toxic-environmental causes. The morphological patterns involved are not of a monotonous kind, but vary according to the cause or agent, the affected site and the gestational age when the abnormality takes effect. This review is concerned with NMD that affects the neocortex, the cerebellum and the brainstem. The neural crest and its disorders will not be included because extensive reviews on this topic have appeared elsewhere.

THE PROCESS OF NEURONAL MIGRATION IN THE BRAIN

Neocortex

The ventricular and subventricular zones of the telencephalon provide the neuronal and glial stem cells and from here migration to the cortical plate, the future neocortex, starts in a radial centrifugal fashion. The migration of young neurons is guided from an early stage by a system of radial glial fibers that span the width of the thickening telencephalon. In the human fetus this process takes place for the greater part between 7 and 16 weeks gestational age. The perikarya of the radial glial cells are in the ventricular and subventricular zones. Cells of glial and neuronal lineage (the former marked by GFAP-staining) could be separated as different proliferative lines within the ventricular epithelium in monkey fetus. The layers of the neocortex are generally laid down in an inside-out fashion, e.g. layer III neurons arriving before layer II neurons which means that later migration waves have to pass earlier migration...
waves. As an exception to this rule it has been suggested that neurons of layer I, the giant Cajal-Retzius neurons and layer VIb, the lower part of layer VI are laid down as a single neuronal network, the primordial plexiform layer, in analogy to the amphibian neocortex and prior to the other layers in mammals. This primordial plexiform layer is thought to provide a framework for the successive migration waves as these become sandwiched between the upper and the lower part of this structure.

The radial glial system was described early in this century by Ramón y Cajal, who used the Golgi technique. Revival of this technique combined with transmission electron microscopy and the application of GFAP-staining facilitated the discovery and detailed description in the monkey fetus. By the use of the same methods demonstration was possible in the human fetus. Choi and Lapham noted the persistence of radial glia in the human telencephalon beyond 16 weeks, raising the possibility that migration does not end sharply at this time. The date of the final plate starts.

Differentiation into neuronal classes characteristic of each cortical layer follows on the completion of migration. This differentiation effects both the shape of the perikaryon (pyramidal and non-pyramidal) and its connections. The commitment of a neuron to differentiate into a certain morphological class appears to depend mostly on the order in which it is generated, rather than on its final position within the neocortex. The best available evidence is the murine reeler mutant that harbors an inherited NMD which specifically affects the intracortical part of the migration trajectory for pyramidal neurons. In the case of this mutant the perikaryal shapes characteristic of each layer are established in spite of faulty positioning. After the completion of the migration process the radial glia disappears. In part these cells appear to transform into astrocytes and ependymal cells. Evidence obtained from the study of fetal human spinal cord suggests that transformation into oligodendroglial cells is also a possibility. One structure belonging to the emerging human neocortex, the subpial layer of Ranke, still awaits elucidation. It occupies the superior part of the molecular layer in the form of several layers of apparent germinal cells from the end of the fourth fetal month in the human until the end of gestation. It was described by Ranke who credits His with the discovery. The structure is only transiently present in some gyrencephalic mammals including man.

The cerebellum

In the cerebellum the mode of migration is different for Purkinje cells and granule cells. The former migrate at 9-10 weeks, but the precise mode is unknown. The granule cells (and possibly the stellate- and basket cells as well) are derived from the external granule cell layer, emerging at 10-11 weeks from the edges of the rhombencephalic roof near the lateral recess of the fourth ventricle, a place where the proximity of the ventricular zone, the starting place of neural cell generation, to the pial surface is very close. From here the future external granule cells start to cover the whole cerebellar surface under the pia. From here postmitotic external granule cells migrate inward and pass the Purkinje cells to form the internal granular layer. In doing so they leave a neurite in the molecular layer that grows out to form the parallel fibers. In this way the external granular layer (and the corpus pontobulbare to be described below) is an exception to the pattern of migration that usually proceeds from the center of the neuraxis in a centrifugal fashion. The guidance for the migrating granule cells in the cerebellum is provided by the vertically oriented Bergmann glial fibers that override the molecular layer with endfeet at the pial surface. The glial cerebellar guidance system has been confirmed in the human fetus. The external granular layer is the latest germinal layer in the brain to disappear, involution starting at 9 months postpartum.

Pontine and olivary nuclei

An important role is played here by the corpus pontobulbare a transient structure near the lateral recess of the fourth ventricle. It represents an accumulation of dividing stem cells located ventrally and anteriorly to the lateral recess where the distance between the ventricular epithelium and the pia is minimal. From here postmitotic neurons destined for the olivary-, arcuate- and pontine nuclei migrate to their final destinations. This represents another instance of a superficially located germinal center. The original site of the cells giving rise to the olivary nuclei is relevant to the location of olivary heterotopia to be described below.

New developments

Research on the process of neuronal migration at the molecular level has only begun. Logic requires that cell-cell recognition especially neuron-glia recognition forms an essential part of the story. Much has been gained already from the study of the autosomal recessive murine weaver mutant. In homozygous weaver granule cells in the cerebellum completely fail to reach the internal granule layer and Bergmann glial cells are severely deficient. Large numbers of arrested granule cells die. Heterozygous weaver shows mildly disturbed migration and abnormal Bergmann glial cells with thickened and irregular processes. To disentangle the respective role of granule cells and Bergmann glial cells chimaceras were produced carrying both heterozygous weaver and normal cell lines that could be distinguished by an enzymatic histochemical marker. In this study it was shown that granule cells carrying the weaver gene were unable to migrate even in the presence of Bergmann cells, whereas the genetically normal granule cells migrated normally. In another study dissociated cultures of weaver cerebellar cells showed both glial and granule cells to be abnormal compared to controls, the former showing stunted growth and the latter dying prematurely. The culture study further showed the existence of two types of astroglia, an extended type resembling Bergmann-glia and a stellate type reminiscent of the internal granular layer. Agglutination studies with a number of lectins demonstrated abnormal surface properties of the cerebellar cells of the weaver mutant. Present evidence recently reviewed suggests that the trophic influence of neurons and astroglia is bidirectional. The experience with the weaver mutant highlights a relation between NMD and cell death that may have significance for the understanding of human pathology. The coincidence of microencephaly and NMD in a high proportion of human cases might in part be explained by similar mechanisms. A recently discovered class of issue specific glycoproteins called cell adhesion molecules...
(CAM), some of them transiently present on the surface of embryonic cells, are now being explored for their role in embryonic shaping processes including neuronal migration. Special interest is focused on a CAM that promote heterotypic (different cell type) adhesion between neuron and glial cell so-called Ng-CAM (neuron-gliala), which has been isolated from chick brain. Beside CAM substrate adhesion molecules (SAM) have been studied intensively. These molecules that differ from CAM include laminin, fibronectin and type IV procollagen. These molecules play a key role in the migration of embryonic cells outside the neuraxis such as neural crest cells but their role within the neuraxis has yet to be decided. The migration of cell processes (neurites) prior to the formation of synapses may have some relevance to the migration of whole cell bodies in terms of the process of cell-cell recognition. Studies with invertebrate species of grasshopper and drosophila have proved the existence of highly specific recognition markers on neuronal cell bodies that provide cues to the exploring growth cone and filopodia of an outgrowing neurite.

Another interesting field possibly related to neuronal migration is polyamine metabolism. Polyamines are low-molecular-weight amines, called spermine and spermidine and their precursor putrescine. These ubiquitous compounds are intimately linked to DNA synthesis and probably also to the synthesis of microtubules and microfilaments. A potent inhibitor of polyamine synthesis, α-difluoromethylornithine (DFMO) exists. When administered to rats between postnatal days 1-21 cerebellar hypoplasia results, combined with entrapment of migrating cerebellar granule cells in the molecular layer.

Clinical experience has focused attention on the possible roles of peroxisomal and mitochondrial fatty acid oxidation in the genesis of inherited NMD such as present in Zellweger (cerebro-hepato-renal) syndrome and warty dysplasia with the genesis of inherited NMD such as present in Zellweger (cerebro-hepato-renal) syndrome and warty dysplasia with (cerebro-hepato-renal) syndrome and warty dysplasia with (cerebro-hepato-renal) syndrome and warty dysplasia with (cerebro-hepato-renal) syndrome and warty dysplasia. 

### Classification of Neuronal Migration Disorders by Morphology

A general classification of NMD is presented in Table 1. A relatively large number of cases with NMD harbors more as one type listed in Table 1. Some types of NMD listed are often seen together e.g. type I agyria and olivary heterotopia or warty dysplasia and leptomeningal neural tissue collections. It is believed that this classification will help the reader to orient himself in the large spectrum of NMD encountered in clinical practice.

**Agyria/pachygyria**

Agyria, otherwise called lissencephaly denotes a smooth brain without secondary sulci. Pachygyria, a related condition, denotes a brain with a thickened neocortex and paucity of secondary sulci. Combinations of the two occur within the same brain. In purely descriptive terms two major types of agyria have been defined as well as a number of case reports that await definite classification.

The first is an order type of migration arrest called classic lissencephaly. It is represented by an abnormal neocortex consisting of the components of the layers III, V and VI combined, separated by a cell sparse zone from radially aligned rows of non-migrated neurons, that often extend to the subependymal zone (Figure 1). The four layered sequence thus defined consists of layer 1, corresponding to the molecular layer, layer 2 harboring neurons with the morphology of the normal layers III, V and VI, layer 3 which is cell sparse and layer 4 which contains heterotopic neurons. Other features regularly seen are decreased brain size, leading to microcephaly, widened ventricles representing a fetal stage in development rather than hydrocephalus and an uncovered Sylvian fossa, representing failure of opercularization. Together with the thickened cortex these macroscopic features allow detection of agyria/pachygyria by neuroradiological means (Figure 2). Additional microscopic features are olivary heterotopia, lodged anywhere between the corpus pontobulbare and their normal station, cerebellar granule cell heterotopia and abnormally shaped dentate nuclei. Aberrant lateral corticospinal tracts in the spinal cord have been described. Purely pachygryic brain may lack accompanying olivary heterotopia. Beside various visceral and other malformations that may be associated a peculiar facial dysmorphia distinguishes a number of reported cases with classical lissencephaly. The phenotype consists of a high forehead, hollow temples, receding chin and vertical wrinkling of the forehead when crying. It has been identified as the eponym, the Miller-Dieker syndrome. Familial occurrence was suggested by unbanded karyograms. Further studies using high resolution chromosome banding revealed anomalies involving the terminal segment of chromosome 17p (one ring chromosome 17p) and one unbalanced translocation resulting in monosomy 17p13) by Dobyns et al. Further studies using high resolution chromosome banding revealed anomalies involving the terminal segment of chromosome 17p (one ring chromosome 17p13) by Dobyns et al. Further studies using high resolution chromosome banding revealed anomalies involving the terminal segment of chromosome 17p (one ring chromosome 17p13) by Dobyns et al. Further studies using high resolution chromosome banding revealed anomalies involving the terminal segment of chromosome 17p (one ring chromosome 17p13) by Dobyns et al. Further studies using high resolution chromosome banding revealed anomalies involving the terminal segment of chromosome 17p (one ring chromosome 17p13) by Dobyns et al. Further studies using high resolution chromosome banding revealed anomalies involving the terminal segment of chromosome 17p (one ring chromosome 17p13) by Dobyns et al. Further studies using high resolution chromosome b
lissencephaly revealed by autopsy was reported by Norman et al.\textsuperscript{53} High resolution banding applied to the parents' karyogram was later reported to be normal.\textsuperscript{51} The patients' facial features related in another paper\textsuperscript{54} are different from Miller-Dieker syndrome and another eponym, the Norman-Roberts syndrome, was proposed to classify the finding. According to a proposal by Dobyns et al.\textsuperscript{54} syndromes representing classical lissencephaly are to be called type I lissencephaly. Macroscopical features allowing recognition by CT-scanning have been defined\textsuperscript{55} (Figure 2).

The second major type of lissencephaly was first described by Walker in 1942.\textsuperscript{56} It is characterized by an almost total disorder of cortical layer formation. Instead of horizontal layers the neocortex is represented by clusters and columns of neurons perpendicular to the surface (Figure 1). This type of lissencephaly has been documented as part of an autosomal recessive disorder under the mnemonic HARD ± E syndrome which stands for Hydrocephalus - Agyria - Retinal dysplasia with or without Encephalocele by Pagon et al.\textsuperscript{57} The eye anomalies that form part of this syndrome affect both the anterior and posterior segments. More or less regular features include microphthalmia (often one-sided), Peter's anomaly, angle...
anomalies, cataracts, persistent hyperplastic vitreous as well as retinal detachment, retinal dysplasia and optic nerve hypoplasia. Characteristic features of the brain include gliomesenchymal proliferation in the leptomeninges encroaching on the underlying neocortex forming septa and investing the mesencephalon. The cerebral cortex — just as in clinical lissencephaly — is not the only part of the brain affected by NMD. The cerebellar folia are fused and a severe layering disorder with Purkinje-cells and granule cells lying haphazardly are seen in every case of the syndrome. In addition, the cerebellum is hypoplastic with absence of the posterior vermis and a Dandy-Walker cyst. Another feature is hypoplasia of the ventral pons with severe reduction of its nuclei and a seemingly hyperplastic corpus pontobulbare. As distinct from classical lissencephaly the inferior olivary nucleus is in its usual place without heterotopic remnants. Encephalocele or occipital dermal sinus are occasionally seen. Another characteristic feature of this disorder concerns white matter abnormalities with paucity of axons and oligodendroglia and severe hypomyelination. The corpus callosum and septum are often absent. Hydrocephalus is usually present, and probably related to the leptomeningeal abnormalities affecting CSF-flow. Since Warburg was the first to draw attention to the genetic syndrome comprising retinal dysplasia and hydrocephalus her contribution was recognized by the proposed name Warburg syndrome, instead of the mnemonic HARD = E. Others suggested calling it Walker-Warburg syndrome (WWS) in regard of Walker's original contribution. A number of papers have served to delineate the clinical, genetic and pathological features. The cerebral pathology of WWS is reminiscent, though not identical, to another autosomal recessive syndrome called Fukuyama's congenital cerebromuscular dystrophy (F-CMD). This syndrome is mainly though not exclusively seen in Japan. The muscular pathology is similar to congenital muscular dystrophy without cerebral involvement. The pathology of the brain is characterized by microgyria with patches of agyria, the latter mainly in the temporal lobes, gliomesenchymal proliferation obliterating the subarachnoid space. Sparserness of myelinated axons, fused frontal poles and cerebellar cortical dysplasias and heterotopia are regularly seen. The disorder is inherited as an autosomal recessive trait and the patients may survive into adulthood, though severely handicapped. A survey of 24 Japanese autopsy cases lists 15 with partial agyria or pachgyria, 3 cases with cataract (unilateral in one patient). In one case all the features beside the muscular dystrophy were characteristic of WWS including retinal detachment and occipital dermal sinus. Descriptions of the neocortical dysplasia especially the agyric regions resemble WWS. This applies both to the mesenchymal obliteration of the subarachnoid spaces and the nodular arrangement of the neocortical neurons seen in WWS. Description of muscle pathology is sparse in WWS. Normal muscle was described in one report. I found changes in various muscles of a personal case of WWS consistent with congenital muscular dystrophy (unpublished). A Dutch sibship has been neuropathologically studied with the main findings common to both WWS and F-CMD. Beside congenital muscular dystrophy and cerebral findings characteristic of F-CMD and WWS the proband had eye anomalies. The latter anomalies not previously published consisted of persistent pupillary membrane, persistent hyaloid artery, small whitish optic nerve heads and pigment layer abnormalities. Other cases have been described that apparently compound WWS and congenital muscular dystrophy. One feature shared by WWS and F-CMD is hypo- or dysmyelination which may be of considerable help in the diagnosis of both conditions. It is not yet certain whether the two conditions represent alleles of one autosomal recessive gene. The combined occurrence of rather unique features such as the rare type of neocortical dysplasia with gliomesenchymal proliferation, cerebellar cortical dysplasia and congenital muscular dystrophy may argue in favor of a single gene involved for both conditions with WWS representing a more severe and earlier onset of the disruptive dysembryonic process. The type of lissencephaly belonging to WWS has been proposed as type II lissencephaly by Dobyns et al.

Besides type I and type II lissencephaly other syndromes with lissencephaly await further studies. One is the exceedingly rare autosomal-recessive Neu-Laxova syndrome a lethal neonatal disease with extreme microencephaly and lissencephaly as well as grotesque skin abnormalities with ichthyosis, collodium-skin and subcutaneous edema. Extreme neopallial hypoplasia with agyria and almost vestigial cerebellum has been described in siblings. The neocortex was represented in the best preserved places by two layers separated by a layer without recognizable neurons. The arrangement bore similarity to type I lissencephaly with the upper neuronal layer probably representing the true cortex and the lowest layer probably representing heterotopic (non-migrated) neurons, but the intralaminar disarray compared to type I lissencephaly was greater. Other cases with microencephaly, lissencephaly and severe underdevelopment of derivatives from the rhombic lips have been described. The designation "cerebro-cerebellar lissencephaly" has been proposed by Dobyns, although the homogeneity of this group remains to be established. For a comprehensive review on less established lissencephaly syndromes the reader is referred to recent papers. As a general feature it is interesting to note that the classical lissencephaly as well as cerebro-cerebellar lissencephaly are not only disorders of neuronal migration but also disorders of organ size.

**Microgyria, brain warts and nodular heterotopia**

Microgyria refers to small meandering gyri without intervening sulci or with intervening sulci apparently bridged by the fusion of the overlying molecular layers (Figure 3). Microgyria

![Figure 3](https://www.cambridge.org/core)
is synonymous with polymicrogyria or micropolygyria, but should be distinguished from sclerotic microgyria or ulogyria, a pure encephaloclastic lesion resulting in atrophic small gyri and relatively broad intervening sulci. Early contributions and theories concerning origin were discussed by Bielschowsky. The histological features of microgyria are not uniform. Layering abnormalities are the rule and mostly of two different kinds: four-layered and unlayered. The four-layered type shows the sequence: marginal layer (top) — neuronal layer — cell-sparse layer with astrocytes— neuronal layer. In the unlayered type no cell-sparse zone is seen dividing upper and lower neuronal strata. A principal cause of microgyria is a circulatory disorder in utero. Frank destruction presenting as full-thickness cavities of the cerebral hemispheres (porencephalies in the classical sense of that term) are often surrounded by areas of microgyria. Microgyric regions in turn will be continuous with normal neocortical areas. Other causes of microgyria are genetic, chromosomal, infectious and toxic, to be discussed below. The mechanism that causes microgyria has not been fully settled. In one case-analysis it was concluded that the lower cell-sparse layer of the four-layered type represented neuronal loss and glial replacement similar to laminar cortical necrosis in the adult with hypoxic-ischemic cortical necrosis and that it represented ipso facto a post-migration accident. The theory was backed up by Golgi analysis of neuronal subtypes in the microgyric cortex which indicated that neuronal classes present in the neocortex — apart from in situ inversions — were in their proper positions.

Czech investigators produced local microgyria in newborn rats by coagulation of the upper half of the neocortex. In this animal neuronal migration to the neocortex is still ongoing at term birth and is completed by the fourth postnatal day. Lesions made on the fourth day failed to produce microgyria. If partial necrosis, e.g. necrosis with the vascular bed preserved was induced in the upper half of the developing cortex young neurons that arrived after the lesion would migrate through the zone of partial necrosis and settle on top of this zone in a disordered way. If necrosis was of sufficient depth a microsulcus would be produced similar to human microgyria. These elegant experiments led these investigators to explain the cell-sparse zone in four-layered microgyria as the result of necrosis and the upper cellular zone as distorted migration after the accident when appropriate guidance by radial glia has been lost because these fibers did not escape necrosis.

Human fetal pathology is sparsely blessed with experiments by nature that provoke microgyria by a single accident of short duration. Two dated carbon monoxide accidents to pregnant mothers at 20 to 24 weeks and 24 weeks gestational age caused four-layered microgyria in the surviving fetuses. (This is far beyond the time at which proliferation of neuroblasts destined for the neocortex grossly ends: 16 weeks, but it may be kept in mind that considerable numbers of young neurons generated before that time still continue to migrate afterward.) In another case the dating of an accident causing microgyria was provided by parabiotic twins, of which one died in utero and the other after fullterm birth. In these monochorionic twins an accident, presumably feto-fetal transfusion, caused death in one and vascular brain damage with survival in the other. The longest survivor of the two had local microgyria (overlying nodular periventricular heterotopias) in a vascular distribution. Dating of the catastrophe was provided mainly by x-ray analysis.

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Another cause of microgyria is intrauterine infection, particularly cytomegaly. Indirect evidence suggests that microgyria is not the result of direct viral attack but results from general perfusion failure. The extent of microgyria is quite variable from case to case. While severe cases may show signs of neurodevelopmental delay and often microcephaly, a mild microgyria restricted to limited neocortical areas may be associated with milder deficiency. A particular case recorded by Galaburda, et al. was that of a man with developmental dyslexia, mild learning disorder and epilepsy. A number of genetic or probably genetic disorders are known to produce microgyria such as Meckel-Gruber syndrome, thanatophoric dysplasia, Fukuyama’s cerebromuscular dystrophy, Bloch-Sulzberger syndrome. Microgyria also occurs in two well defined inherited disorders of metabolism related to peroxisomai dysfunction. In one of these, Zellweger’s cerebro-hepato-renal syndrome, results in periventricular, subcortical and intracortical heterotopia. The neocortex is often referred to as both microgyric and pachygyric, but differs from both these conditions. The microgyric aspect in Zellweger syndrome is apparently the result of fusion of distinct small gyri. The four layer pattern is not found and microgyri also line the bottom of sulci, a phenomenon referred to as “cloverleaf microgyria”. Regions that appear macroscopically pachygyric in Zellweger syndrome are histologically almost similar to the microgyric regions. Both neurons and glial cells show light microscopic and ultrastructural changes in Zellweger syndrome and an impressive storage of lipid material of various types is seen in macrophages and astrocytes. The mechanism of NMD in Zellweger syndrome is yet unknown. Since a number of metabolic pathways are involved, all resulting from the absence or near-absence of peroxisomes no deficit can be singled out as the cause of NMD in this complex disorder. Another related autosomal recessive disorder called neonatal adrenoleukodystrophy (NALD) has deficient peroxisomes and NMD expressed as areas of microgyria in addition to sudanophilic leukodystrophy and adrenal atrophy. The genetic relationship to Zellweger syndrome has still to be ascertained in depth at this time.

Brainwarts (verruccous dysplasia, dysgénésie nodulaire de l’écorce) Brainwarts present microscopically as tiny “ herniations” of the second neocortical layer into the first layer, thereby reaching the surface (Figure 4). To the naked eye the abnormality presents as a flat, round, often dimpled disk seated on the crown of a gyrus, less often in the depth of a sulcus. The phenomenon was first described in 1873 by Simon and called brain wart by Jakob in 1940.
frontal lobes and the Rolandic areas. Mild warty dysplasia has a remarkably high incidence varying between 16% and 26% of routine autopsies if carefully looked for. A common origin with microgyria has been suggested. In another study it has been shown that not just the upper layers but all the cortical layers may participate in the formation of the “wart”. An apparently related phenomenon, often seen in autopsies of immature fetuses up to 24 weeks presents microscopically as fountains of cortical neurons apparently bursting into the first layer, that is still smooth (agyric) in accordance with fetal age. Larroche believes that it represents a pathological phenomenon (status verrucosus simplex) related to microgyria. Specific associations with verrucose dysplasia are rare. One that deserves mention is neonatal glutaric aciduria type II or multiple acyl-CoA dehydrogenase deficiency, a disorder that affects mitochondrial beta-oxidation. The association with verrucose dysplasia has been described in male sibs. The neocortical dysgene­sis consisted of symmetrical reduction of the number of gyri of frontal, parietal and temporal lobes and an irregular surface with numerous warty protrusions. Microscopically these “warts” consisted of multiple small gyri that were partially fused as well as heterotopic neuronal clusters in the molecular layer and the subcortical white matter. In addition bile duct hypoplasia, cholestasis, siderosis and fatty degeneration were found in the liver of both infants as well as enlarged bilateral polycystic kidneys.

**Leptomeningeal heterotopias**

Heterotopic collections of astrocytes with or without admixture of ectopic neurons are often observed in conjunction with heterotopic invasions of the first neocortical layer. It appears that such heterotopia are provoked by discontinuities in the external limiting membrane that is made up by glial endfeet. Leptomeningeal heterotopia may be seen together with verrucose dysplasia (Figure 4). Large glo-neuronal heterotopia in the leptomeninges have been described in cases of familial microencephaly, pachygyria and congenital nephrotic syndrome. Leptomeningeal heterotopia are not rare. They may be seen in cases of holoprosencephaly, environmental causes of NMD (to be described below) and in vascular disruptions. Leptomeningeal glial heterotopia may be seen surrounding the brainstem e.g. the mesencephalon in cases of Walker-Warburg syndrome. Experimentally leptomeningeal heterotopia have been provoked in neonatal rat by application of the drug 6-hydroxydopamine which causes a breach in the barrier of glial endfeet formed in the cerebellum by Bergmann glial cells as well as the basal lamina. These breaches caused the appearance of external granule cells in the subarachnoid space between the folia as well as fusion of adjacent folia.

**Nodular neuronal heterotopia in the cerebral hemispheres**

Heterotopic neuronal masses represent the clearest example of NMD (Figure 5). These can occur anywhere along the migration trajectory. In the telencephalon they may occur mostly in the subependymal zone or just below the neocortex. Their degree of cytological maturation varies and may be quite impressive, to the extent that pyramidal and nonpyramidal neurons may be distinguished and both subtypes may carry abundant numbers of well developed dendritic spines in Golgi sections. The maturation achieved is likely to result in biological activity of a false kind because of improper wiring due to ectopic positioning. Large heterotopic clusters are not likely to arise after the main bulk of migrating neurons has arrived at the cortical plate, that is after the 16th week of gestation. The causes of nodular heterotopia are extremely varied and include genetic, chromosomal, vascular and environmental causes. These various causes are therefore described in the appropriate sections. The size of such heterotopia is usually small, often below the resolution afforded by CT- or MRI-scanning apparatus. Sizable masses may occasionally be picked up by either means (Figures 6, 7). Subependymal heterotopia may cause bulging of the ventricular wall, but this is not a reliable sign unless absorption characteristics (CT) or better T1 weighted MRI images suggest grey matter.

**Schizencephaly and allied disorders**

Connatal clefts in the brain mantle may be accompanied by NMD. Full thickness defects that cause continuity between the arachnoid space and the lateral ventricles have been called porencephalies by Heschl (1859). With respect to the areas surrounding such defects these may exhibit: (1) destruction of the adjacent neocortex and white matter without NMD (2) microgyric neocortex with the histopathological structure of the four-layered type (3) neocortical and heterotopic collect-

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**Figure 4** — Verrucous cortical dysplasia in 6 weeks old premature (35 weeks) born infant with multiple congenital anomalies, with normal karyogram. Undiagnosed syndrome. HE. 95x.

**Figure 5** — Periventricular heterotopic nodular masses in a newborn with occipital encephalocele. HE. 12.7x.
tions adjacent to the deeper part of the cleft up to the ventricular wall. Yakovlev was the first to describe the third category under the name schizencephaly. He distinguished two types of schizencephaly. In the first type the lips of the cleft were apposed by a so-called pia-ependymal seam. In the second type the lips of the cleft were open. The latter type was associated with hydrocephalus. Clefts were covered with ectopic grey matter. Yakovlev considered schizencephaly a true malformation. As such it has become a classic subdivision amongst fetal neurodevelopmental disorders.

It remains difficult, however, to follow Yakovlev’s conception of such defects as a type of focal malformation. The absence of inflammatory or gliotic lesions noted by Yakovlev does not exclude an extrinsic origin since this absence is usual in early fetal disruptions. On the other hand no familial cases or cases associated with chromosomal disorders have been reported that would support a programming failure (true malformation) as the cause of schizencephaly. It may therefore be reasonable to consider Yakovlev’s schizencephaly and Heschl’s porencephaly with full thickness defect parts of a spectrum of fetal disruptions. At one end are the post-migration period accidents resulting in lesions without associated NMD. On the other hand fetal vascular damage might be suspected as the cause of schizencephaly. It remains difficult, however, to follow Yakovlev’s conception of such defects as a type of focal malformation. The absence of inflammatory or gliotic lesions noted by Yakovlev does not exclude an extrinsic origin since this absence is usual in early fetal disruptions. On the other hand no familial cases or cases associated with chromosomal disorders have been reported that would support a programming failure (true malformation) as the cause of schizencephaly. It may therefore be reasonable to consider Yakovlev’s schizencephaly and Heschl’s porencephaly with full thickness defect parts of a spectrum of fetal disruptions. At one end are the post-migration period accidents resulting in lesions without associated NMD. On the other hand fetal vascular damage might be suspected as the cause of schizencephaly.

Cerebellar cortical dysplasias and heterotopia

The usual type of heterotopia in the cerebellum is a sharply defined patch containing granule cells, molecular layer and Purkinje cells apparently thrown together in a more or less haphazard way. Small collections of this type or containing only granule cells as the neuronal component may be seen postmortem in normal infants, mainly in the floccular and nodular lobes. Gross lesions of this composition, macroscopically visible within the white matter or continuous with normal adjacent cerebellar cortex represent more serious malformations. In the case of continuity of the heterotopic cortex with the normal cortex the name cerebellar (poly)microgyria has been given. Because no excessive folding or small gyri are involved (as in the case of the cerebral counterpart) Friede has absence of the septum pellucidum and blunted lateral ventricular angles due to heterotopia. These features as well as occasionally the heterotopia rimming the cleft may be discovered by CT-scanning (Figure 6).

Olivary heterotopia

Heterotopia involving olivary components anywhere between the corpus ponto-bulbare and their normal station are a regular feature of the Miller-Dieker syndrome. Similar lesions have been occasionally in cases with Dandy-Walker syndrome without other distinguishing features, and on one occasion in Coffin-Siris syndrome (Table II).
Table 2: Syndromes with neuronal migration disorders

<table>
<thead>
<tr>
<th>Classification</th>
<th>Genetics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Metabolic syndromes</td>
<td>ar, xr</td>
</tr>
<tr>
<td>Zellweger s.</td>
<td>ar</td>
</tr>
<tr>
<td>Neonatal adrenoleukodystrophy</td>
<td>ar</td>
</tr>
<tr>
<td>Glutaric aciduria II</td>
<td>ar</td>
</tr>
<tr>
<td>Menkes' disease</td>
<td>ar</td>
</tr>
<tr>
<td>G1M gangliosidosis</td>
<td>ar</td>
</tr>
<tr>
<td>Neuromuscular syndromes</td>
<td>ad</td>
</tr>
<tr>
<td>Walker-Warburg s.</td>
<td>ad</td>
</tr>
<tr>
<td>Fukuyama syndrome</td>
<td>ad</td>
</tr>
<tr>
<td>Myotonic dystrophy</td>
<td>ad</td>
</tr>
<tr>
<td>Anterior horn arthropathy</td>
<td>ad</td>
</tr>
<tr>
<td>Neurocutaneous syndromes</td>
<td>xd</td>
</tr>
<tr>
<td>Incontinentia pigmeni</td>
<td>xd</td>
</tr>
<tr>
<td>Neurofibromatosis</td>
<td>ad</td>
</tr>
<tr>
<td>Ito's hypomelanosis</td>
<td>xd</td>
</tr>
<tr>
<td>Encephalocraniocutaneous lipomatosis</td>
<td>xd</td>
</tr>
<tr>
<td>Tuberous sclerosis</td>
<td>xd</td>
</tr>
<tr>
<td>Epidermal nevus s. (Jadassohn)</td>
<td>xd</td>
</tr>
<tr>
<td>Multiple congenital anomalies-syndromes</td>
<td>?</td>
</tr>
<tr>
<td>Smith-Lemli-Opitz s.</td>
<td>?</td>
</tr>
<tr>
<td>Oligohydramnios tetrad (Potter s.)</td>
<td>?</td>
</tr>
<tr>
<td>Cornelia de Lange s.</td>
<td>?</td>
</tr>
<tr>
<td>Meckel-Gruber s.</td>
<td>?</td>
</tr>
<tr>
<td>Oro-facio-digital s.</td>
<td>?</td>
</tr>
<tr>
<td>Coffin-Siris s.</td>
<td>?</td>
</tr>
<tr>
<td>Chromosomal syndromes</td>
<td>ad</td>
</tr>
<tr>
<td>Trisomy 13</td>
<td>ad</td>
</tr>
<tr>
<td>Trisomy 18</td>
<td>ad</td>
</tr>
<tr>
<td>Deletion 4p</td>
<td>ad</td>
</tr>
<tr>
<td>Deletion 17p13 (Miller-Dieker s.)</td>
<td>ad</td>
</tr>
<tr>
<td>Skeletal dysplasias</td>
<td>xd</td>
</tr>
<tr>
<td>Thanatophoric dysplasia</td>
<td>xd</td>
</tr>
<tr>
<td>Nephrotic syndrome</td>
<td>ad</td>
</tr>
<tr>
<td>Pachygyria/nephr. s. (Robain)</td>
<td>ad</td>
</tr>
<tr>
<td>Other CNS-dysplasias</td>
<td>ad</td>
</tr>
<tr>
<td>Aicardi s.</td>
<td>ad</td>
</tr>
<tr>
<td>Joubert s.</td>
<td>ad</td>
</tr>
<tr>
<td>Type I lissencephaly, normal karyot. (Norman- Roberts)</td>
<td>ad</td>
</tr>
<tr>
<td>Cerebro-cerebellar lissencephalies</td>
<td>ad</td>
</tr>
<tr>
<td>Hemimegalephalies</td>
<td>ad</td>
</tr>
<tr>
<td>Schizencephaly and allied s.</td>
<td>ad</td>
</tr>
<tr>
<td>Twin-syndromes</td>
<td>?</td>
</tr>
<tr>
<td>Parabiotic twin syndrome (early)</td>
<td>?</td>
</tr>
<tr>
<td>Solitary reports</td>
<td>?</td>
</tr>
</tbody>
</table>

Abbreviations: ar = autosomal recessive; ad = autosomal dominant; xd = x-linked dominant; xr = x-linked recessive.

Table 3: Maternal and environmental causes of neuronal migration disorder.

<table>
<thead>
<tr>
<th>Infection</th>
<th>Intoxications</th>
<th>Ionizing radiation*</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>cytomegalovirus</em></td>
<td><em>carbonmonoxide</em></td>
<td><em>ibid.</em></td>
</tr>
<tr>
<td><em>carbonmonoxide</em></td>
<td><em>isotretinoic acid</em></td>
<td><em>ibid.</em></td>
</tr>
<tr>
<td><em>ethanol</em></td>
<td><em>methylmercury</em></td>
<td><em>ibid.</em></td>
</tr>
<tr>
<td><em>isotretinoin</em></td>
<td></td>
<td></td>
</tr>
<tr>
<td><em>ionizing radiation</em></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Limited evidence in man, but high probability in view of animal experiments; see text.

NMD. The first two are outside the scope of this article. Dysgenesis of the corpus callosum is found relatively often in infants with grossly disturbed mental development and epilepsy especially infantile spasms. The association between dysgenesis of the corpus callosum and NMD — both microgyria and nodular neuronal heterotopia — is so close that it is found irrespective of etiology. It is therefore very likely that similar mechanisms underlie both NMD and callosal dysgenesis. In most cases of callosal dysgenesis the origin of the corpus callosum is not absent, but represented by paired ectopic longitudinal bundles of Probst. A aberrant neurite outgrowth is therefore an essential feature of callosal dysgenesis.

The interrelation with aberrantly placed perikarya, the essence of NMD, is a tempting area for future research. Among the rare but specific causes of this association the Aicardi syndrome should be mentioned. A comprehensive recent review is available. In this syndrome which is exclusively present in the female sex, or at least in individuals having two X-chromosomes, choriotretinal lacunae, dysgenesis of the corpus callosum, vertebral anomalies and clinical patterns of severe developmental retardation and infantile spasms are found. Neuropathological studies reviewed mention cortical lamination disturbance as well as subcortical and subependymal heterotopia. Other supratentorial brain abnormalities reported in Aicardi syndrome are "porencephaly", hemispheric cysts and anomalies of the choroid plexus including papilloma. In at least one autopsy case an interhemispheric neuroepithelial cyst was found. This suggests that some of the cysts seen on CT-scans of patients with Aicardi syndrome may be similar neuroepithelial cysts. Such cysts are believed to result from dislodged ventricular epithelium early in development. The association of callosal dysgenesis, NMD and neuroepithelial cysts may therefore be of more than incidental significance.

In another syndrome that is probably X-linked dominant, the oral-facial-digital syndrome, NMD is found together with callosal dysgenesis and occasionally neuroepithelial cysts. In one report congenital coloboma in one retina and hypoplastic optic nerves were found as well, providing some interesting parallels with Aicardi syndrome. A relationship that may exist between NMD and intraparenchymal neuroepithelial cysts has been found in experimental animals (rats), subjected to prenatal irradiation. The cysts originate from "neuroblast" rosettes.

Chromosomal disorders

Severe mental deficiency is expressed in most of the known chromosomal disorders. This predicts a high association with structural brain defects. Unfortunately the harvest of neuropathological observations has been small compared with the huge body of literature dealing with these disorders. Even where abnormalities have been found such as dysgenesis of the corpus callosum, such findings often did not explain the severity of neurological handicap. The elucidation of this problem had to await a more subtle technique such as the revival of the Golgi staining technique that revealed the abnormalities of the synaptic organisation of the neocortical neurons e.g. in trisomy 13 and in Down's syndrome. Gross abnormalities such as holoprosencephaly in trisomy 13 and myelomeningocele in trisomy 18 are well known. NMD in trisomy 13 usually takes the form of heterotopic collections in the cerebellar white matter. Many cases of trisomy 18 show periventricular heterotopia in the cerebral hemispheres.
heterotopias are also known in the 4p- syndrome, beside abnormalities of gyration, microgyria, increased numbers of neurons in the molecular layer of the neocortex and Parkinjie cell heterotopias. Trisomy 21 (Down syndrome) is well known for a combination of developmental and regressive abnormalities. Occasional mention has been made of nodular heterotopias in the cerebral white matter and mixed heterotopias of variable size in the cerebellar flocculus.

Multiple congenital anomalies (MCA-syndromes) and NMD

Beside chromosomal syndromes hereditary or genetically undetermined MCA-syndromes may carry a high incidence of NMD. Some of these have already been mentioned. In the genetic group autosomal recessive disorders include the Smith-Lemli-Opitz syndrome with heterotopia in the cerebral and the cerebellar hemispheres, especially in those cases in which the full syndrome, including polydactyly and renal polycystic disease is expressed. Cerebellar heterotopias have been described in infants with Potter syndrome (oligohydramnios tetrad), and combined cerebral and cerebellar heterotopia in Cornelia de Lange syndrome. Furthermore NMD is seen in Meckel-Gruber syndrome (autosomal recessive), Zellweger syndrome, glutaric aciduria type II with brain warts and renal cysts. The oro-facio-digital syndrome has been mentioned already. Besides the typical features of facial skull, extremities and cerebral malformation it may also feature polycystic kidneys. The association between NMD and renal dysplasias in otherwise widely different MCA-syndromes may be significant.

NMD associated with neurocutaneous syndromes

von Recklinghausen neurofibromatosis is associated with frank mental retardation in a small number of cases. An autopsy study of patients with this phacomatosis revealed mild abnormalities in cortical architecture, especially in those whose intelligence was subnormal. Gross malformation consisting of microgyria and nodular heterotopias was observed in a case with IQ 39.

Tuberous sclerosis, the second neurocutaneous syndrome is particularly associated with mental retardation in a high proportion. Important abnormalities found in autopsied patients include disturbances of glial differentiation and growth of a topical nature, including subventricular nodules and giant cell tumors. Although tuberous sclerosis has a well documented prenatal onset in many cases reported, NMD does not appear a significant part of the morphological abnormalities encountered. One report describes malpositioning of pyramidal neurons in a cortical tuber studied with the Golgi technique. Of the rarer neurocutaneous syndromes grey matter heterotopias together with glial proliferation has been observed in Ito’s hypomelanosis. Microgyria has been reported in encephalocranio-cutaneous lipomatosis and leptomeningeal glioneural and white matter heterotopias, together with microgyria and gliomatosis in a newborn with severe epidermal nevus syndrome. Microgyria in Bloch-Sulzberger syndrome has already been mentioned.

NMD associated with hemimegalencephaly

A number of pathological case reports exist on infants and young children with hemimegalencephaly, a condition with one hyperplastic cerebral hemisphere with gyral abnormalities (pachygryria), giant pyramidal neurons (restricted to the pathological side), beside subcortical and glioneural leptomeningeal heterotopias. Cytomorphometric studies in some of these cases proved increased nuclear volume and an apparently increased DNA content in the affected neurons, which led to a suggestion of topical heteroploidy. The presence of giant neurons in the brainstem ipsilateral to the giant hemisphere in some cases and the presence of ipsilateral corporeal hypertrophy in some cases (reference list of Bignami, et al 1968) would imply that the dysembryonic influence causing this growth disturbance is rather limited to one side of the main embryonic axis, and therefore may well originate during the earliest mitotic divisions of the embryo.

The presence of glial nodules and giant glial cells in the absence of gross degenerative changes in some of the reported cases is reminiscent of disorders affecting growth and proliferation in a topical nature, in other words the phacomatoses. The latter opinion concurs with the pathological findings in an autopsy case of the one neurocutaneous syndrome that causes hemimegalencephaly: the organoid nevus syndrome or epidermal nevus syndrome. The hemimegalencephaly cases are also remarkable for they present rare examples of brain malformations with NMD in which brain volume is increased rather than decreased. An MRI-example of hemimegalencephaly is shown in Figure 7.

NMD associated with megalencephaly and elevated insulin-like growth factor II

A single case report on congenital megalencephaly with grossly disturbed neocortical development and NMD with elevated levels of the growth hormone dependent insulin like growth factor II (IGF II) in CSF (at autopsy) and in postmortem brain samples appeared recently. This interesting study offers a new approach to cases of “intrinsic” disturbances of bulk growth whether associated with NMD or not.

Figure 7 — Hemimegalencephaly in a 2 year old female demonstrated in transverse inversion recovery sequence MRI-section. The abnormal hemisphere seen on the right shows paucity of secondary sulci, deep parietal sulcus and masses of poorly delineated grey matter within the central white matter.
Confirmed hazards to neuronal migration in the human fetus are isotretinoic acid, ethanol, methylmercury, radiation and radiomimetics. The effects of fetal hypoxia have already been mentioned.

**Isotretinoic acid**

Isotretinoic acid, an alcohol-soluble synthetic analogue of vitamin A prescribed as an oral medication for severe cystic acne has become associated with craniofacial, cardiac, thymic and central nervous system malformations in fetuses exposed during the first trimester. A spectrum of cerebral abnormalities have been described which includes hydrocephalus, microcephaly, holoprosencephaly (one case), vermal aplasia, cerebellar cortical dysplasia, dysorphic corticospinal tracts in the brainstem, malformed inferior olivary nucleus, malformed allocortex, focal neocortical agyria. A consistent abnormality appears leptomeningeal neuroglial heterotopia that may affect both supra- and infratentorial structures. 

**Ethanol**

In utero exposure to ethanol produces the fetal alcohol syndrome (FAS), a dysmorphic syndrome with in utero as well as postnatal growth retardation, a characteristic facial dysmorphia with prominent midfacial hypoplasia, microcephaly, mental retardation and often cardiac defects. Increased rate of stillbirth is another recognized hazard. Morphological brain abnormalities are variable and logically depend on time and degree of exposure and possibly on additional adverse conditions such as dietary deficiencies and other addictions including heavy smoking. A spectrum of neuropathological findings has been reported in infants and fetuses, which includes microencephaly, hydrocephalus, arhinencephaly, callosal hypogenesis microdysplasias of cerebral and cerebellar cortices, dentate- and olivary nuclei, hydromyelia, porencephaly and spongy degeneration in diencephalic structures and optic nerves. NMD is mainly seen as leptomeningeal neuroglial heterotopia of various extent overlying both supra- and infratentorial parts of the neuraxis. Such neuroglial heterotopia appear to arise through thin bridges of neural tissue that connect the heterotopia with the underlying neuraxial structures. Neuronal heterotopia within the cerebral hemispheres are occasionally found. The neuropathological series quoted undoubtedly are the most serious part of the spectrum of sequelae. Moderate mental retardation and microcephaly with behavioral disorders, characteristically present in long-term survivors may have other structural correlates than NMD. A large number of animal experiments involving different species and different protocols of exposure all point to the potentially damaging effects of ethanol on the shaping process in various parts of the brain.

**Methylmercury**

In the 1950’s methylmercury was the cause of large scale industrial pollution around Minamata Bay (Japan) carried by consumption of poisoned fish from the bay. So-called Minamata disease caused severe neurological deficits. Also babies who were exposed in utero were affected by fetal Minamata disease. Severe neuronal losses in the cerebral and cerebellar cortices were described, but also signs of NMD. Another epidemic of methylmercury intoxication in Iraq (1970-1971) was caused by consumption of homemade bread prepared from seed grain of wheat treated with methylmercury fungicide. Prematally exposed babies suffered from psychomotor retardation even when the clinical symptoms in their mothers had been mild or absent. The brains of two infants expiring soon after birth have been described in detail by Choi et al. They had been exposed between 6 and 8 and between 8 and 10 weeks fetal age. Mercury intoxication was confirmed by its determination in the blood and was found to be higher in the infants at delivery as in their mothers, confirming delayed fetal clearance. The babies were small for gestational age. Major findings consisted of neuronal heterotopia in the cerebral hemispheres and in the cerebellum, and leptomeningeal heterotopia. In the cerebral cortex, layering abnormalities and undulating upper cortical layers resembling microgyria were apparent. Large numbers of gemistocytic astrocytes containing mercury were shown histochemically. A relatively large number of reports relate to the influence of methylmercury on experimental animal fetuses (for review see Choi 1983). The influence of methylmercury on migrating neurons has been studied in vitro by exposing human fetal explants containing migrating neurons to methylmercury. Methylmercury chloride caused abrupt cessation of active movement of cells in these cultures. The initial site of damage appeared to be the neuritic membrane in the vicinity of growth cones.

Electron microscopy suggested that the initial event was the disappearance of neurotubules necessary for structural support and for axoplasmic transport. Similar damage was observed in cultures of astroglial cells. Decreased DNA-synthesis probably resulted from interference with mitotic spindles. The outcome of these studies may have a bearing not only on public health policies surrounding organic mercury, but also on other agents and intrinsic processes that affect cytoskeletal proteins and in this way affect neuronal migration.

**Ionizing radiation and cytostatic drugs**

Pregnant rats subjected to roentgen irradiation between the 14th and 16th day produce offspring with neuronal heterotopias in the cerebral hemispheres. If radiation is applied between the 19th and 21st day disordered cerebellar migration is found. Other deficiencies observed were microencephaly, absent corpus callosum, hydrocephalus and rosette formations. Cerebellar granule cell ectopia were seen in rats after birth if radiation took place before migration from the external granule cell layer ended. Similar observations could be made with respect to the cerebral hemispheres in mice when irradiated between the 10th and 14th day of gestation.

The experience in man has been summarized by the description of the sequelae in survivors of prenatal exposure to the atomic bomb of Hiroshima and by the timetable of the effects of prenatal radiation injury obtained from a number of case-reports on therapeutic pelvic irradiation during pregnancy. In the case of the Hiroshima bomb microcephaly (below – 2 S.D.) was the most obvious sequel and this was especially prevalent in cases that had been exposed between the 7th and 15th week gestational age. Most though not all of the cases had learning disorders. Later analyses have confirmed this. Fetal exposure to pelvic radiation, mainly due to the vague of radiation for all kinds of purposes in the twenties and thirties has provided another source. A timetable constructed from these individual reports showed that radiation incurred between 3 to 4 and 11 weeks caused microcephaly, mental retardation.
and stunted growth, besides eye, skeletal, and genital abnormalities. Between 11 and 16 weeks radiation resulted in microcephaly, mental retardation and stunted growth without associated injury. Similar effects though milder were encountered in cases of radiation between 16 and 20 weeks. Cerebellar NMD has been described in a case where intraperitoneal radium had been applied ending "near the seventh month".201 Through the sparsity of detailed neuropathological studies the results of experimental studies cited above find no confirmation or exclusion in man. Since the effects that can be observed in man during life such as eye abnormalities, stunted growth and microcephaly are closely similar to those that are encountered in animal experiments the likelihood of NMD being eventually found in surviving human cases through the use of magnetic resonance imaging or postmortem investigation is high. Animal experiments with cystostatic drugs e.g. cytosine arabinoside found in surviving human cases through the use of magnetic resonance imaging or postmortem investigation is high. Animal experiments with cystostatic drugs e.g. cytosine arabinoside indicate similar results as those obtained with ionizing radiation.202203

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REFERENCES


118. Jacob H. Die feinere Oberflächengestaltung der Hirnwindung: die Hirnwarzenbildung und die Mikropolygyrie. Z. Gesamte Neurol Psychiatr 1940; 170: 64-68.


208. Moerman Ph, Barth PG. Olio-petto-cerebellar atrophy with muscular atrophy, and joint contractures of prenatal onset. Virchow's Arch A (in press).
