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Three tachykinins in mammalian brain

Two polypeptide precursors to the neuromodulatory substance P have recently been identified. One of them (β-preprotachykinin) contains amino acid sequences corresponding not only to substance P but also to substance K, a novel, related peptide. A third substance P-like peptide, neuromedin K, has recently been isolated from spinal cord. The existence in vertebrates of three members of the tachykinin family of peptides may account for pharmacological observations suggesting the presence of more than one type of substance P receptor in the nervous system.

The tachykinin family of peptides (Fig. 1) is characterized by the common amino acid sequence - Phe-X-Gly-Leu-Met-NH₂ at the C terminus, where X is a hydrophobic or aromatic residue. Substance P, the best-known member of the family, was until recently the only tachykinin known to occur in mammalian tissues other than amphibians. However, a number of lines of evidence suggested that there might be other tachykinins present in vertebrate tissue.

Using a variety of antisera to the amphibian peptide physalaemin, Lazarus and colleagues were able to demonstrate the widespread occurrence in mammals of material with physalaemin-like immunoreactivity. Immunological, chemical and pharmacological analysis of the physalaemin-like material in a small-cell carcinoma from human lung demonstrated that it was similar or identical in its properties to amphibian physalaemin. Keen and colleagues demonstrated the biosynthesis in dorsal root ganglia of substance P together with a related peptide which was immunoprecipitated by C-terminal but not by N-terminal substance P antisera. Analysis of the incorporation of different radiolabelled amino acids into the novel peptide indicated that it differed from substance P in containing no proline residues and in containing the amino acids phenylalanine and methionine in the ratio 1/1 instead of 2/1.

Evidence for the existence of novel tachykinins in mammalian tissue was complemented by evidence for the presence of more than one type of tachykinin receptor in vertebrates. This concept, first suggested by Erspamer, was extended by Iversen and colleagues who defined the responses of a variety of pharmacological test systems in terms of two types of tachykinin receptor. All the tachykinins display approximately equal potency at one type of receptor (the SP-P receptor), whereas the second type of receptor (the SP-E receptor) is considerably more sensitive to elephakin and kassinin than to the other tachykinins. Searching for an endogenous ligand to the SP-E receptor, Maggio et al. developed a radioimmunoassay for kassinin, the most potent ligand at SP-E receptors. They found that extracts of spinal cord contained abundant kassinin-like immunoreactivity which they attributed to a novel peptide. This peptide they named 'substance K' to indicate its apparent immunochemical similarity to kassinin.

Kimura et al. were at this time searching for novel neuropeptides in porcine spinal cord, using as an assay their activity on the isolated guinea-pig ileum. Four fractions exhibiting ileum-contracting activity were identified by Sephadex G-15 chromatography: two of the fractions contained substance P and its oxidized derivative, the remaining fractions contained a novel peptide.
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destined for secretion from the cell, the α- and β-PPT sequences begin with a 15–30 amino acid ‘signal sequence’ (largely composed of hydrophobic amino acids), which promotes attachment of the newly synthesized polypeptide to the endoplasmic reticulum and its secretion into the intracisternal space.

This sequence is cleaved off by a membrane-bound protease during or immediately after synthesis: the exact length of the signal sequence in the PPT molecule has not yet been determined. Immediately preceding the sequence of substance K in the precursor is a pair of basic amino acids (Lys-Arg). A trypsin-like protease is thought to cleave the precursor after the second basic residue to liberate the N terminus of the substance K molecule. The N terminus of the substance P sequence (Arg-Pro) is preceded in the precursor by a single basic amino acid (arginine). Although the double basic amino acid ‘signal’ for proteolytic processing is again present, the Arg-Pro bond is selectively resistant to trypsin-like proteases and instead the cleavage occurs between the two arginine residues. Immediately following the sequences of substance P and substance K is the sequence -Gly-Lys-Arg — a pair of basic amino acids indicating a site of proteolytic processing, preceded by a glycine residue which is thought to be the donor of the amide group found at the C terminus of both peptides.

Other biologically active peptides may be liberated from the PPT sequences. In particular the sequence -Arg-Arg-Lys at the C terminus of both precursors could be a processing site, liberating peptides corresponding to residues 72–108 of α-PPT and residues 111–126 of β-PPT. These peptides may possess some, as yet undefined, biological activity.

The presence of two precursors for substance P in brain has exciting implications. The two PPT molecules may arise from two closely similar but separate genes. Alternatively they may arise from a single gene by alternative splicing events. The vast majority of eukaryotic genes consist of precisely defined regions coding for sections of mature mRNA (exons) interspersed with regions (introns) which are either deleted during the synthesis of mRNA precursors or ‘spliced’ out of the mRNA sequence during its maturation. The portion of the β-PPT mRNA that is missing from α-PPT may correspond to a single exon, and both precursor mRNAs may be generated from a common gene product by different splicing events. A similar variable splicing mechanism enables a single gene (the calcitonin gene) to code for different products (calcitonin and calcitonin-related peptide) in the thyroid and hypothalamus, respectively. The origin of the two PPT molecules will finally be understood (no doubt, very soon) when the structure of the PPT gene(s) is established.

Regardless of their origin, the existence of two tachykinin precursors in brain poses intriguing problems for neurobiology. Substance P-containing neurons may express either α- or β-PPT molecules throughout their lifetime, implying the presence of separate groups of ‘substance P only’ and ‘substance P + substance K’ neurons in the brain. Alternatively a given neuron may be able to switch its pattern of peptide production between the two states either during development or in response to external stimuli. In this context, it may be particularly worthwhile to examine the effects of nerve growth factor in stimulating substance P synthesis in sensory neurons, and the effects of decentralization in increasing depolarization in decreasing – SP synthesis in autonomic ganglia. Are the relative amounts of substance P and substance K synthesized also affected?

These questions will doubtless be resolved soon: the production of specific antisera to substance K (and indeed to neuropekinin K) will permit the distribution of these peptides in the brain, and any differences between them, to be established. cDNA probes to those parts of the PPT mRNA common to both precursors, and to the region missing from α-PPT, will be used to identify in-situ hybridization those cells producing each of the two tachykinin mRNAs and to look for plastic and developmental changes in their levels in brain.

This leaves neuropekinin K as something of a Cinderella in the substance P family, as we still know nothing of its distribution, or biosynthetic origin in the brain. It is attractive to imagine that this peptide, which like substance K is probably a good ligand at SP-E receptors, may exist in a new set of peptidergic neurons in brain. The combined distribution of ‘substance P only’, ‘substance P + substance K’ and ‘neuropekinin K only’ neurons might then account for the observed regional differences in substance P receptor subtypes in the nervous system.

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these data and the absence of a consistent abnormality in all patients with MS, a combination of influences and factors has been postulated. The most common hypothesis is that the patient with MS is genetically susceptible in some manner, possibly by way of an immunogenetic abnormality related to the major histocompatibility complex, and that a particular environmental exposure, presumably through an infectious agent such as a virus, leads to the destruction of CNS myelin. However, explanations for recurrences, unpredictable progression and marked variations in intensity of the disease have not yet been approached by these theories.

There are few chronic neurological diseases of humans which may be convincingly demonstrated to result from virus infection. These include progressive multifocal leukoencephalopathy due to a papovavirus, subacute sclerosing panencephalitis resulting from rubella, progressive rubella panencephalitis, and several post-infectious encephalomyelitides. In none of these is the pathology exactly like that of MS, nor have any of these agents been identified in MS tissue. In fact the most current techniques of virus isolation or recognition of virus genomic material have not shown positive or specific results in MS tissue. It is true that a number of claims have been made about isolation of viruses from MS tissue, but in no case have these claims been confirmed.

Animals afford a more controllable situation in which to study CNS myelin injury, and several diseases have been popular models for studies of both viral causes and autoimmunity as they might relate to MS. One of these is experimental allergic encephalomyelitis (EAE), an autoimmune response to a component of CNS myelin called myelin basic protein, injected with complete Freund’s adjuvant which appears to be an essential but not understood facilitator. Myelin basic protein, which has a monomeric molecular weight of 18 500 and comprises approximately 30% of CNS myelin proteins, has been extensively studied because of this role in provoking EAE. Susceptibility to EAE varies among different animal species and strains. In rodents, the Lewis rat is the most susceptible. Cellular immunity against myelin basic protein plays the primary role in the immunopathogenesis of acute EAE. Thus, EAE may be passively transferred to uninunized recipients with mononuclear cells from lymphoid tissue of sensitized animals.

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models which permit the induction of a strain of mouse hepatitis virus. In particular, the acute form of EAE differs from MS in being more intense, usually monophasic, and pathologically showing more CNS inflammation than demyelination. However, there are now several disease models which permit the induction of a subacute or chronic form of EAE which resembles the clinical course of MS more closely, exhibiting periods of exacerbation and remission, and which also exhibits pathology more similar to that of MS.'2,13 At least in the guinea-pig, the age of the animal is important for the development of chronic, as opposed to acute, EAE; juvenile guinea-pigs are the most susceptible to induction of the chronic form.

Viral diseases of animals which have been studied as possible models of MS include visna in sheep, canine distemper, Thielers murine encephalomyelitis, and CNS infections produced by the JHM strain of mouse hepatitis virus. In particular, the latter two examples have recently been studied in depth because they offer models which allow close scrutiny. Thielers murine encephalomyelitis virus, a picornavirus, infects oligodendrocytes which then bear the virus antigen and serve as targets for a host immune response against the affected cells.14 The JHM strain of mouse hepatitis virus is a coronavirus which selectively destroys oligodendrocytes in its natural host, the mouse.15,16 The CNS effects of the JHM strain have recently been given a fresh look by ter Meuwen and co-workers, who have provided evidence of subacute demyelination in a fraction of 4-5-week-old Lewis rats inoculated intracranially with the virus.17 A prominent feature of this demyelination was a perivascular mononuclear cell infiltration which resembled in some respects the pathology of EAE. Following this lead, it was then shown that lymphocytes from these animals had been sensitized to myelin basic protein. Furthermore, if these lymphocytes were exposed to myelin basic protein in vitro, they acquired the ability to transfer subacute demyelination passively into recipients. Control studies indicate that virus or myelin basic protein alone is neither being transferred nor causative in this passively transferred disease. Thus, this model provides preliminary information that a protracted virus infection of an unnatural and immature host may lead to subsequent demyelination and sensitization to a myelin component.

The availability of this model should be very useful in further assessing the events intervening between a virus infection and the appearance of inflammatory demyelination. Whether this will provide incisive clues as to the etiology of MS remains to be seen, but at least it ties in with the possibility that the age of the animal or other temporal events influence the intensity and course of the disease may provide important new insights into the cause of MS.

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