

Paraneoplastic Encephalomyelitis Opinion and Treatment Encephalitis is a Seditious Condition of the Brain with Numerous Etiologies

Abstract

The neuroendocrine system has close interactions with the immune system. Their bidirectional communications

head of he fe p hing on he cer i ignal he relea e of o ocin from he po erior pi i ar of he mo her. O ocin hen ra el o he er here i im la e erine con rac ion . e elici ed erine con rac ion ill hen im la e he relea e of increa ing amo n of o ocin. i po i i e feedback loop ill con in e n il par ri ion. Since e ogeno l admini ered and endogeno l ecre ed o ocin re l in he ame e ec on he female reprod c i e em, n he ic o ocin ma be ed in peci c in ance d ring he an epar m and po par m period o ind ce or impro e erine con rac ion .

Ceo ocin ne on-imm ne ne o k

em i mainl compo ed of e o ocin- ecre ing magnocell lar o ocin ne ron in he praop ic n cle, para en ric lar (PVN) n clei and e eral h po halamic acce or n clei, he po erior pi i ar harboring heir a onal erminal, heir a ocia ed glial cell and pre nap ic ne ron ha direc l reg la e o ocin ne ron ac i i ie. e par ocell lar para en ric lar o ocin ne ron are ano her branch of he o ocin- ecre ing em and he major o rce of brain and pinal cord o ocin, hich ha e clo e in erac ion i h he magnocell lar o ocin ne ron [4]. In hi em, o ocin ne ron can en e change in nap ic inner a ion , a roc ic ac i i , blood-borne fac or , and elf-relea ed chemical ell a he le el of imm ne c okine in he local ne ral circ i . а 0 ocin ne ron b eq en l in egra e he e ignal and reg la e

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I pla a i al role in labo r and deli er . e hormone i prod ced in he h po halam and i ecre ed from he para en ric lar n cle o he po erior pi i ar here i i ored. I i hen relea ed in p l e d ring childbir h o ind ce erine con rac ion . e concen ra ion of o ocin recep or on he m ome ri m increa e igni can l d ring pregnanc and reache a peak in earl labor. Ac i a ion of o ocin recep or on he m ome ri m rigger a do n ream ca cade ha lead o increa ed in racell lar calci m in erine m o bril hich reng hen and increa e he freq enc of erine con rac ion . In h man, mo hormone are reg la ed b nega i e feedback; ho e er, o ocin i one of he fe ha i reg la ed b po i i e feedback. e imm nologic ac i i ie b relea ing o ocin in o he blood and he brain. Corre pondingl, o ocin recep or (OXTR) are e en i el e pre ed in cen ral and peripheral i e incl ding cla ical imm ne organ, i e and cell, ch a monoc e and macrophage, h mic T-cell, and me ench mal romal cell of ad l bone marro . , o ocin can mod la e ac i i ie of bo h he inna e and acq ired imm ne em hile e er ing broad e ec on he ac i i of cen ral and peripheral i e .Con er el, o ocin ne ron al oe pre man c okine recep or, ch a in erle kin (IL)-6 and recei e mod la ion of imm ne em form af nc ional ni in o r bod ' defen e em.

Rela i e o o ocin, he kno n a ocia ion of a opre in i h imm ne em i mo l indirec and limi ed. e imm nologic reg la or role of a opre in i likel d e o i promo ion of adrenocor ico ropin hormone relea e . In m rine h m , OTXR pre en in all T-cell b e , m ch broader han he pre ence of a opre in recep or . Ne ralizing o ocin b no a opre in ing peci c an ibodie ind ce a marked increa e in IL-6 and le kemia inhibi or fac or ecre ion in cell c l re . Rela i e o he clear imm nologic e ec ha blocking OXTR igni can l inhibi he prod c ion of c okine IL-1 and IL-6 elici ed b an i-CD3 rea men of h man hole blood cell c l re , he imm nologic f nc ion of a opre in ere largel no eri ed. , e en a i el belie e ha he o ocin-b no a opre in- ecre ing em i he major carrier in ne roendocrine reg la ion of imm nologic ac i i ie ia he ne roh poph i [5].

Pa ici a ion of he o ocin- ec e ing em in la e ed imm nologic defen e

e imm ne em pro ec he bod again di ea e hro gh de ec ing pa hogen, pre en ing heir in a ion/di ion, red cing heir inj r e ec and eradica ing hem from he bod . e o ocinecre ing em e ec e he e f nc ion hro gh hree la ered defen e i h increa ing peci ci ha incl de he rface barrier, he inna e and he adap i e imm ne proce e .

S rface barrier : e mo primar form of imm ne defen e em i he rface barrier ha incl de he ph ical and chemical barrier . e ph ical barrier can pre en pa hogen ch a bac eria and ir e from en ering he organi m. A prereq i i e of e ec ing hi f nc ion i he r c ral in egri of he barrier like he kin, blood-brain barrier, and in e inal m c a ell a indi id al cell and i e.O ocin in ol e hi la er of defen e a r b i an ibio ic abili and o nd-healing e ec [6]. I ha been repor ed ha in pa ien i h diabe e melli, o ocin inhibi he focal micro ora of p o-in amma or proce e and lead o a more rapid elimina ion of microorgani m from he p o-in amma or foc . Moreo er, local applica ion of o ocin increa e he e cac of cipro o acin in rea ing ep ic o nd . ro gh enhancing he f nc ion of cla ical an ibio ic and direc an imicrobial e ec, o ocin can accelera e o nd clo re b promo ing a c logene i and prolifera ion of endo helioc e and hi ioc e , and h increa e kin re i ance o pa hogen infec ion . a locall applied o ocin promo e he barrier f nc ion i al o a ocia ed i h i an i ecre or and an i lcer e ec . S bc aneo applica ion of o ocin canno onl red ce b rnind ced kin damage b al o alle ia e ga ric and ileal in amma ion and damage b red cing i e ne rophil in l ra ion and TNFrelea e. Moreo er, o ocin can reng hen he in e ine m co a barrier b ind cing pro aglandin E2 relea e. In addi ion, o ocin can al o main ain he r c ral in egri of cell lar and i e again i chemic inj r a ho n in ra 'kidne , li er , kele al m cle ,

o ar and hear . Similarl , in raperi oneal o ocin admini ra ion accelera e f nc ional, hi ological, and elec roph iological reco er a er di eren cia ic inj r model in ra . B main aining he in egri of indi id al cell , i e and organ em , o ocin can reng hen he ph ical barrier and in rn enhance bod ' defen e abili [7].

Inna e imm ne em: If a pa hogen breache he rface barrier and ge in o he bod, he inna e imm ne em can pro ide an immedia e re pon e b relea ing an ibac erial molec le and mobilizing imm ne cell. Di eren from he ac ion of o her imm nologic mod la or, he e ec of o ocin on he inna e imm ni i a mobilizing he imm ne defen e po en ial hile ppre ing pa hogenic inj r d e o o er-reac ion of he inna e imm ni . A repor ed, o ocin ac on me ench mal romal cell of ad 1 bone marro o promo e bone forma ion and all blood , o ocin can increa e he re er e of imm nologic lineage . capaci . Con er el , lipopol accharide and ep i can increa e pla ma o ocin le el, hich in rn decrea e TNF- and IL-1 le el in he macrophage and red ce pero ide prod c ion in OXTRbearing monoc e and macrophage [8-10]. O ocin al o ppre e endo o in-ind ced increa e in pla ma adrenocor ico ropin hormone TNF-, IL-1, IL-6, and o her c okine. In he an i chemic inj r e ec, o ocin dimini he cell apop o i and bro ic depo i in he remo e m ocardi m hile ppre ing in amma ion b red c ion of ne rophil, macrophage and T-l mphoc e. Al ho gho ocin cold al o e er proin amma or e ec a er , peci call a h man labor, i mainl pla imm nologic homeo a ic role in re pon e o imm nologic challenge. e imm ne-reg la ing f nc ion of o ocin al o pre en in he ran plan a ion of me ench mal em cell. O ocin rea ed mbilical cord deri ed- me ench mal em cell ho a decrea e in be forma ion b a dra ic increa e in ran ell migra ion acii . i e ecia ocia edih he increa ed ran crip ion le el of ma ri me allopro eina e-2 . e o ocin pre rea men al o increa e me ench mal em cell engra men and conne in e pre ion in infarc ed m ocardi m and cardiac con rac ili in ra, hich along i h he inhibi or e ec of o ocin on in amma or c okine relea e o ld facili a e he cce of cell ran plan a ion. Imm node cienc : O ocin can be bene cial o he rea men of h man imm node cienc . For in ance, in ADIS pa ien, he n mber of o ocin ne ron red ce igni can l in he PVN; hro gh increa ing CD4+ cell co n , o ocin can impro e he heal h a of omen infec ed i h HIV.

References

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" problem

bearing rats

5.

bearing rats

rats