

Melatonin

Resolution of: 4 July 2019 Valid until: unlimited

Entry into force on: 4 July 2019

Federal Gazette, BAnz AT 08.08.2019 B4

Therapeutic indication (according to the marketing authorisation of 20 September 2018):

Slenyto is indicated for the treatment of sleep disorders (insomnia) in children and adolescents aged 2–18 years with Autism Spectrum Disorder (ASD) and/or Smith-Magenis syndrome where sleep hygiene measures have been insufficient.

1. Additional benefit of the medicinal product in relation to the appropriate comparator therapy

Children and adolescents aged 2–18 years with sleep disorders associated with Autism Spectrum Disorder (ASD) and/or Smith-Magenis syndrome where sleep hygiene measures have been insufficient.

Appropriate comparator therapy:

Best supportive care.

Best supportive care (BSC) is the therapy that ensures the best possible, patient-individual, supportive treatment to alleviate symptoms and improve the quality of life.

Extent and probability of the additional benefit of melatonin compared with best supportive care:

Hint for a minor additional benefit.

Study results according to endpoints:

Children and adolescents aged 2–18 years with sleep disorders associated with Autism Spectrum Disorder (ASD) and/or Smith-Magenis syndrome where sleep hygiene measures have been insufficient.

Results of the NEU_CH_7911 study 1:

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¹ Data from the dossier evaluation of the IQWiG (A19-04) and from the addendum (A19-47) to the dossier evaluation.

Endpoint category	Melatonin		Placebo		Melatonin vs placebo
Endpoint	N	Patients with event n (%)	N	Patients with event n (%)	RR [95% CI] p value
Mortality					
Overall mortality	60	0 (0)	65	0 (0)	-

N ^a		Endpoint Melatonin category				placebo	
	Values at the start of study MV ^b (SD)	Change at week 15 MV° (SE)	N ^a	Values at the start of study MV ^b (SD)	Change at week 15 MV° (SE)	MD [95% CI] p value ^c	
Morbidity							
no data avail able	457.21 (101.28)	51.03 (10.46)	no data avail able	459.85 (109.22)	18.71 (10.82)	32.32 [2.38; 62.26] 0.035	
no data avail able	95.16 (59.25)	-37.77 (6.82)	no data avail able	98.76 (73.90)	-12.57 (7.01)	-25.20 [-44.61; -5.80] 0.011	
no data avail able	45.5 (19.42)	1.96 (1.33)	no data avail able	47.5 (18.43)	1.84 (1.36)	0.13 [-3.64; 3.89] 0.948	
Behavioural strengths and abnormalities (SDQ)							
no data avail able	20.2 (5.28)	-0.84 (0.39)	no data avail able	21.1 (5.86)	0.17 (0.41)	-1.01 [-2.12; 0.11] 0.077	
no data avail able	3.0 (2.00)	-0.24 (0.14)	no data avail able	3.5 (1.98)	0.05 (0.14)	-0.29 [-0.69; 0.11]	
no data avail able	4.3 (2.69)	-0.11 (0.23)	no data avail able	4.3 (2.98)	-0.02 (0.24)	-0.10 [-0.75; 0.55]	
no data avail able	8.0 (2.00)	-0.47 (0.20)	no data avail able	8.0 (2.27)	0.07 (0.21)	-0.54 [-1.12; 0.03]	
no data avail able	4.9 (2.15)	-0.02 (0.15)	no data avail able	5.4 (2.11)	0.03 (0.16)	-0.05 [-0.49; 0.39]	
No data avail able	4.9 (2.94)	0.21 (0.24)	No data avail able	4.4 (2.89)	0.34 (0.25)	-0.13 [-0.81; 0.55] 0.702	
No data avail able	5.3 (2.84)	-0.57 (0.28)	No data avail able	5.3 (2.69)	0.16 (0.30)	-0.74 [-1.55; 0.08] 0.076	
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Quality of life

No data available.

a: Number of patients included in the evaluation unclear; however, based on the responses it is certain that it is sufficiently

b: Corresponds to the time of randomisation: week 2 of the study
c: Mean and SE (change at week 15 per treatment group) as well as MD, 95% CI, and p value (group comparison): MMRM
d: Higher values mean better function; a positive group difference corresponds to an advantage for melatonin

f: Higher values mean better pro-social behaviour; a positive group difference corresponds to an advantage for melatonin CGAS: Children's Global Assessment Scale; CI: confidence interval; MD: Mean Value Difference; MMRM: mixed model with repeated measurements; MV: Mean Value; N: Number of patients evaluated; RCT: randomised controlled study; SE: standard error; SD: standard deviation; SDQ: Strength and Difficulties Questionnaire; vs: versus

Endpoint category	Melatonin		Placebo		Melatonin vs placebo
Endpoint	N	Patients with event n (%)	N	Patients with event n (%)	RR [95% CI] p valueª
Side effects					
AE (additionally shown) ^b	60	51 (85.0)	65	50 (76.9)	-
SAEs	60	0 (0)	65	1 (1.5)	0.36 [0.01; 8.69] 0.515
Withdrawal because of AE ^b	60	1 (1.7)	65	1 (1.5)	1.08 [0.07; 16.94] > 0.999
Somnolence (PT, AE) ^b	60	17 (28.3)	65	8 (12.3)	2.30 [1.07; 4.94] 0.027

a: IQWiG calculation of RR, 95% CI (asymptotic) and p value (unconditional exact test, CSZ method). In the case of 0 events in one study arm, the correction factor 0.5 was used to calculate RR and CI in both study arms.

2. Number of patients or demarcation of patient groups eligible for treatment

Children and adolescents aged 2–18 years with sleep disorders associated with Autism Spectrum Disorder (ASD) and/or Smith-Magenis syndrome where sleep hygiene measures have been insufficient.

approx. 8,000-86,000 patients

3. Requirements for a quality-assured application

The requirements of the product information are to be taken into account. The European Medicines Agency (EMA) provides the contents of the product information (summary of product characteristics, SmPC) for Slenyto[®] (active ingredient: melatonin) at the following publicly accessible link (last access: 15 May 2019):

https://www.ema.europa.eu/documents/product-information/slenyto-epar-product-information_de.pdf

e: Higher values mean better more problems or a greater impairment; a negative group difference corresponds to an advantage for melatonin

b: Discrepancy between information in Modules 4 A and 5 of the dossier. The data shown are from Module 5.

CI: confidence interval; n: number of patients with (at least 1) event; N: number of patients evaluated; PT: preferred term; RCT: randomised controlled trial; RR: relative risk; SAE: serious adverse event; AE: adverse event; vs: versus

4. Treatment costs

Annual treatment costs:

Children and adolescents aged 2–18 years with sleep disorders associated with Autism Spectrum Disorder (ASD) and/or Smith-Magenis syndrome where sleep hygiene measures have been insufficient.

Designation of the therapy	Annual treatment costs/patient			
Medicinal product to be assessed:				
Melatonin	€928.07-3,744.66			
Best supportive care	different for each individual patient			
Appropriate comparator therapy:				
Best supportive care	different for each individual patient			

Costs after deduction of statutory rebates (LAUER-TAXE®) as last revised: 15 June 2019)

Costs for additionally required SHI services: not applicable