

The Role of the 5-HTTLPR Gene Variation of the SLC6A4 Serotonergic System in the Development of Addictive Disorders: A Narrative Review

Роль вариации 5-HTTLPR гена SLC6A4 серотонинергической системы в формировании аддиктивных расстройств: нарративный обзор литературы

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Review

Alexey Krylov, Nadezhda Pavlova,
Alexey Bochurov

*Yakut Science Centre of Complex Medical Problems,
Yakutsk, Russia*

Алексей Крылов, Надежда Павлова,
Алексей Бочуров

*ФГБНУ «Якутский научный центр комплексных
медицинских проблем», Якутск, Россия*

ABSTRACT

BACKGROUND: Addictive disorders remain a global problem, affecting health, society and the economy. The etiopathogenesis of addictions, which have a multifactorial nature, is poorly understood, making it difficult to develop personalized treatment approaches. Of particular interest is the *SLC6A4* gene, which regulates serotonergic transmission. The 5-HTTLPR variation of this gene is associated with the risk of addictions, but the data are contradictory due to the heterogeneity of clinical manifestations and pleiotropic effects of the gene. Integration of genetic, environmental and neurobiological factors into multidimensional models is becoming relevant.

AIM: The aim of this study is to assess the role of 5-HTTLPR variations in the *SLC6A4* gene of the serotonergic system in the development of addictive disorders.

METHODS: The manuscripts were searched in the MEDLINE and eLIBRARY.RU databases using the keywords in Russian and English: “*SLC6A4*”, “5-HTTLPR”, “addictive disorders”, “pharmacogenetics”, “serotonin”, “antidepressants”, “ethnic differences”. After eliminating duplicates and a two-stage screening (by titles/annotations and full-text analysis) of the 1,561 discovered papers, the final review included 41 publications that meet the stated inclusion criteria.

RESULTS: The S-allele of 5-HTTLPR is associated with an increased risk of addictions and comorbid affective disorders, but its role is ambiguous due to the heterogeneity of symptoms. Ethnic differences have been identified: the S-allele predominates (70.6–80.9%) in Asian populations, the L-allele in Europeans (38.5–66.7%). Unique neurobiological markers for S-allele carriers have not been established, and the pleiotropic effects of *SLC6A4* are also observed in other mental disorders, which reduces its specificity for addictions.

CONCLUSION: The inconsistency of the data on 5-HTTLPR highlights the need to take into account ethnic specificity and develop multivariate models that integrate genetic, environmental and clinical factors. This will improve risk prediction (development of addictions), personalization of therapy and the effectiveness of pharmacogenetic approaches, reducing the likelihood of adverse reactions.

АННОТАЦИЯ

ВВЕДЕНИЕ: Аддиктивные расстройства остаются глобальной проблемой здравоохранения, комплексно влияя на здоровье, социум и экономику. Этиопатогенез зависимостей, имеющих мультифакториальную природу, изучен недостаточно, что затрудняет разработку персонализированных подходов к лечению пациентов. Особый интерес представляет ген *SLC6A4*, регулирующий серотонинергическую передачу. Вариация 5-HTTLPR этого гена ассоциирована с риском развития зависимостей, однако данные противоречивы из-за гетерогенности клинических проявлений и плеiotропных эффектов гена. Актуальной становится интеграция генетических, средовых и нейробиологических факторов в многомерные модели.

ЦЕЛЬ: Оценить роль изменения 5-HTTLPR гена *SLC6A4* серотонинергической системы в формировании аддиктивных расстройств.

МЕТОДЫ: Поиск публикаций производили в базах MEDLINE и eLIBRARY.RU с использованием ключевых слов «*SLC6A4*», «5-HTTLPR», «аддиктивные расстройства», «фармакогенетика», «серотонин», «антидепрессанты», «этнические различия», «*addictive disorders*», «*pharmacogenetics*», «*serotonin*», «*antidepressants*», «*ethnic differences*». После исключения дубликатов и двухэтапного скрининга (по названиям/аннотациям и полнотекстовому анализу) из 1561 обнаруженной работы в финальный обзор вошла 41 публикация, соответствующая заявленным критериям включения.

РЕЗУЛЬТАТЫ: S-аллель 5-HTTLPR ассоциирован с повышенным риском развития зависимостей и коморбидных аффективных нарушений, однако его роль неоднозначна из-за гетерогенности симптомов. Выявлены следующие этнические различия: S-аллель преобладает (70,6–80,9%) в азиатских популяциях, L-аллель — у европейцев (38,5–66,7%). Уникальные нейробиологические маркеры для носителей S-аллеля не установлены, а плеiotропные эффекты *SLC6A4* наблюдаются и при других психических расстройствах, что снижает его специфичность для аддикций.

ЗАКЛЮЧЕНИЕ: Противоречивость данных о 5-HTTLPR подчеркивает необходимость учета этнической специфики и разработки многомерных моделей, объединяющих генетические, средовые и клинические факторы. Это позволит улучшить прогнозирование рисков возникновения аддикций, персонализацию терапии и эффективность фармакогенетических подходов, снижая вероятность развития нежелательных реакций.

Keywords: *5-HTTLPR variant in the SLC6A4 gene; psychogenetics; serotonin; addictive behavior*

Ключевые слова: *изменение 5-HTTLPR гена SLC6A4; психогенетика; серотонин; аддиктивное поведение*

INTRODUCTION

The number of people who are dependent on psychoactive substances is rapidly increasing worldwide, including in Russia [1, 2]. According to the World Health Organization (WHO), alcohol and drug abuse, as well as the use of other psychoactive substances, has become an epidemic in this early 21st century [3, 4]. It should also be noted that the number of affected families by addiction and requiring professional and timely assistance is also on the increase [5]. Differences amongst individuals in the propensity for addictive behavior, including nicotine dependence, are partially mediated by genetic factors [6]. Current estimates

of heritability for all major addictive disorders range from 40% to 80% [7].

Addictive behavior is one form of deviant behavior that arises from the desire to escape reality [8]. The presence of addictive behavior indicates impaired ability to adapt to altered environmental conditions [9]. Addictive behavior traditionally includes alcohol abuse, toxicomania, drug addiction, tobacco smoking (chemical dependencies), as well as computer addiction, gambling, love addictions, sexual addictions, workaholism, and food addiction (overeating, fasting) [10]. Disorders related to psychoactive substance use represent the most common and severe forms of

addiction, classified under the International Classification of Diseases, 10th Revision (ICD-10), code F1: “Mental and behavioral disorders due to psychoactive substance use” [11].

Such functions as mood, emotions, cognition, motor abilities, and circadian and neuroendocrine rhythms — including appetite, sleep, and reproductive activity — are regulated by the serotonin system in the midbrain [12]. Fluctuations in serotonin levels is one of the effects of addictive behavior, underscoring the importance of the genes that encode serotonergic receptors and the transporters in the pathogenesis of dependence [13]. One of the candidate genes that affect the development of dependences is the *SLC6A4* serotonin transporter gene [10]. Recent studies have demonstrated that the 5-HTTLPR (serotonin transporter-linked polymorphic region) variant in this gene is associated with smoking behavior; however, the level of its implication remains inconclusive due to insufficient research [10, 14].

Studies of the 5-HTTLPR pathological allele in the *SLC6A4* gene indicate that there is a connection between various mental disorders and the transcriptional activity levels of the S and L-alleles [15]. For example, reduced activity of the S-allele has been associated with anxiety, depression, suicide attempts, and bipolar disorder, whereas enhanced activity of the L-allele is considered protective against depression, but has also been linked to suicidal behavior, nicotine dependence, and attention-deficit/hyperactivity disorder [15–17]. The aforementioned alleles may also influence treatment efficacy; for example, serotonin reuptake inhibitors may prove more effective in patients with depression and posttraumatic stress disorder who carry the L-alleles [18]. In particular, S-allele carriage is associated with an increased risk of adverse outcomes as relates to alcohol use, mediated by reduced sensitivity to ethanol [19].

The aim of this study is to assess the role of 5-HTTLPR variations in the *SLC6A4* gene of the serotonergic system in the development of addictive disorders.

METHODS

Eligibility criteria

Inclusion criteria:

- original research and meta-analyses regarding the role of the 5-HTTLPR variant in the *SLC6A4* gene in the development of addictive disorders, including interaction of genetic and environmental factors;
- publications analyzing the pharmacogenetic aspects of the use of antidepressants (selective

serotonin reuptake inhibitors, SSRIs) in carriers of different 5-HTTLPR polymorphisms;

- studies related to ethnic differences in the S and L-alleles distribution and their association with clinical outcomes.

Exclusion criteria:

- case reports and case series without the use of control groups;
- publications related solely to therapy for addictive disorders without the analysis of genetic factors;
- publications in languages other than Russian or English.

Information sources

The search was conducted in the electronic databases MEDLINE and eLIBRARY.RU. The search was carried out in December 2024.

The search period covered ran from January 2003 to December 2024. The search was limited to 2003 because that year marked the publication of the first fundamental studies on the role of 5-HTTLPR [20], which laid the foundation for the study of the interaction between this polymorphism and mental disorders, as well as addictive behavior.

Search strategy

The following combination of keywords in Russian and English were used to search for publications: “*SLC6A4*”, “5-HTTLPR”, “addictive disorders”, “pharmacogenetics”, “serotonin”, “antidepressants”, “ethnic differences”. The search for publications was performed in stages. The search sequence is shown in Figure 1.

Selection process

Each publication was identified by a manual search. Several specialists from the group of authors of this article conducted the search and selection of publications (see Authors' contributions section). Some publications selected at the screening stage were excluded from further analysis once it became clear that they did not meet the eligibility criteria (Figure 1).

Analysis of the results

The authors analyzed each publication and summarized information from the selected sources. The results of the summarization are presented in the structured text and figures.

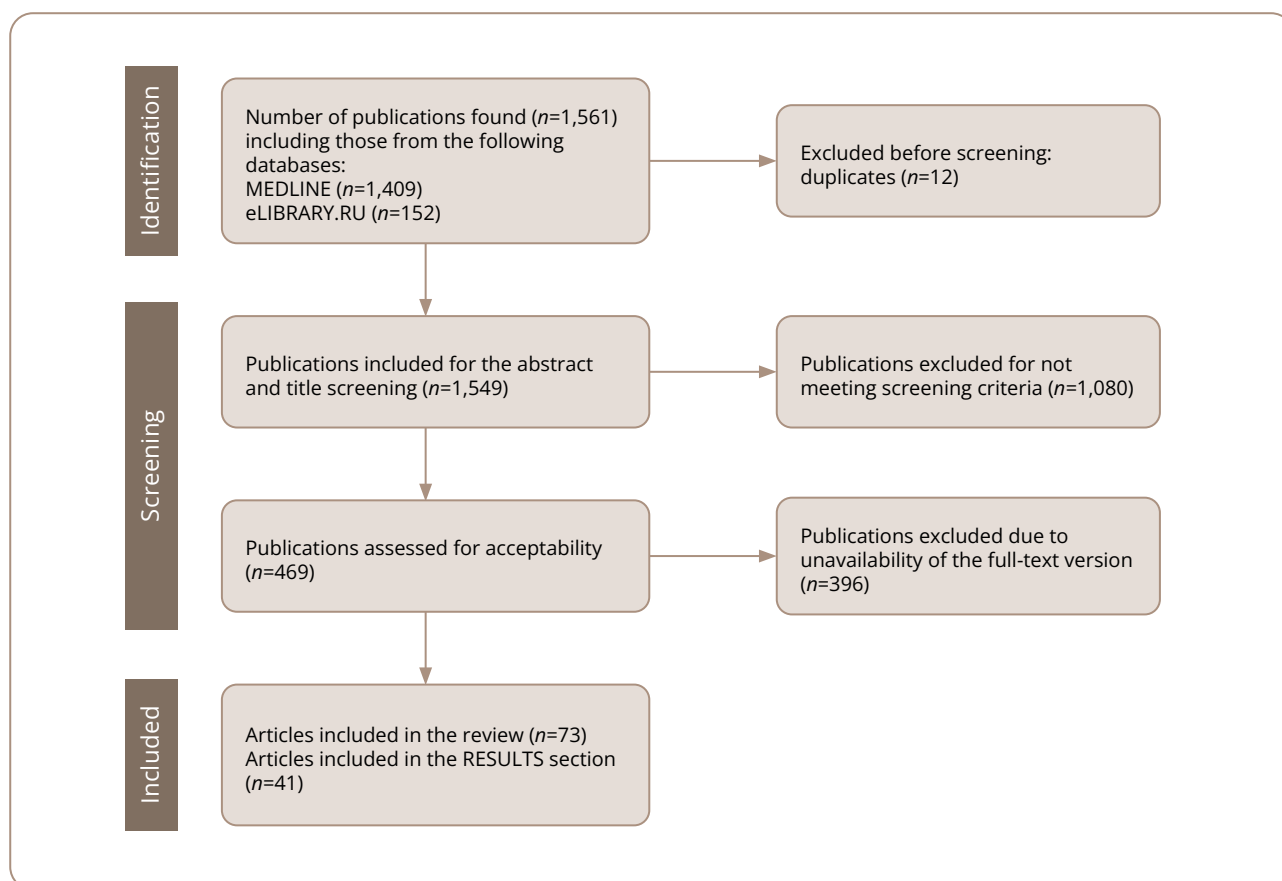


Figure 1. Flow chart demonstrating the selection process.

Source: Krylov et al., 2025.

RESULTS

The *SLC6A4* gene and its relation to psychiatric peculiarities

The 5-HTTLPR (rs4795541) polymorphic region is a functional insertion/deletion polymorphism of 44 base pairs in the promoter region of the *SLC6A4* serotonin transporter gene (Figure 2) [21]. 5-HTTLPR is one of the variants that has most extensively been studied in patients with mental disorders [22–24]. It has also been widely investigated in the context of intermediate phenotypes, such as neuroimaging modalities and gene-environment interactions, with the latter typically examined in relation to affective and anxiety phenotypes [21, 25, 26].

However, it should be noted that addictive disorder is a complicated process whose development depends on a number of factors, including those associated with family history, as well as factors associated with neurobiology, the social context, and experience [14]. This is why the contribution of the pathological allele of the *SLC6A4* gene to mental characteristics may be

only one of many factors influencing this psychological trait [27].

In a study comparing the frequency of pathogenic variations in 5-HTTLPR and rs25531 A<G of the *SLC6A4* gene among the Yakuts and other population samples, a high frequency of the S-allele was identified, which was similar to that observed in Chinese and Japanese populations [14, 25]. According to the study by Nardi et al., the S-allele (deletion) is associated with a lower expression of the serotonin transporter gene [28]. Moreover, carriers of the S-allele demonstrate increased sensitivity to environmental stimuli [28], which likely contributes to the accumulation of this allele among the Yakuts [14].

Some mental disorders with a comprehensive mechanism of pathogenesis (such as schizophrenia) are associated with a disruption of the serotonin system, which affects the development and differentiation of neurons [29]. Moreover, its transporter, encoded by the *SLC6A4* gene, plays a key role in the regulation of the activity level of the serotonergic system [30].

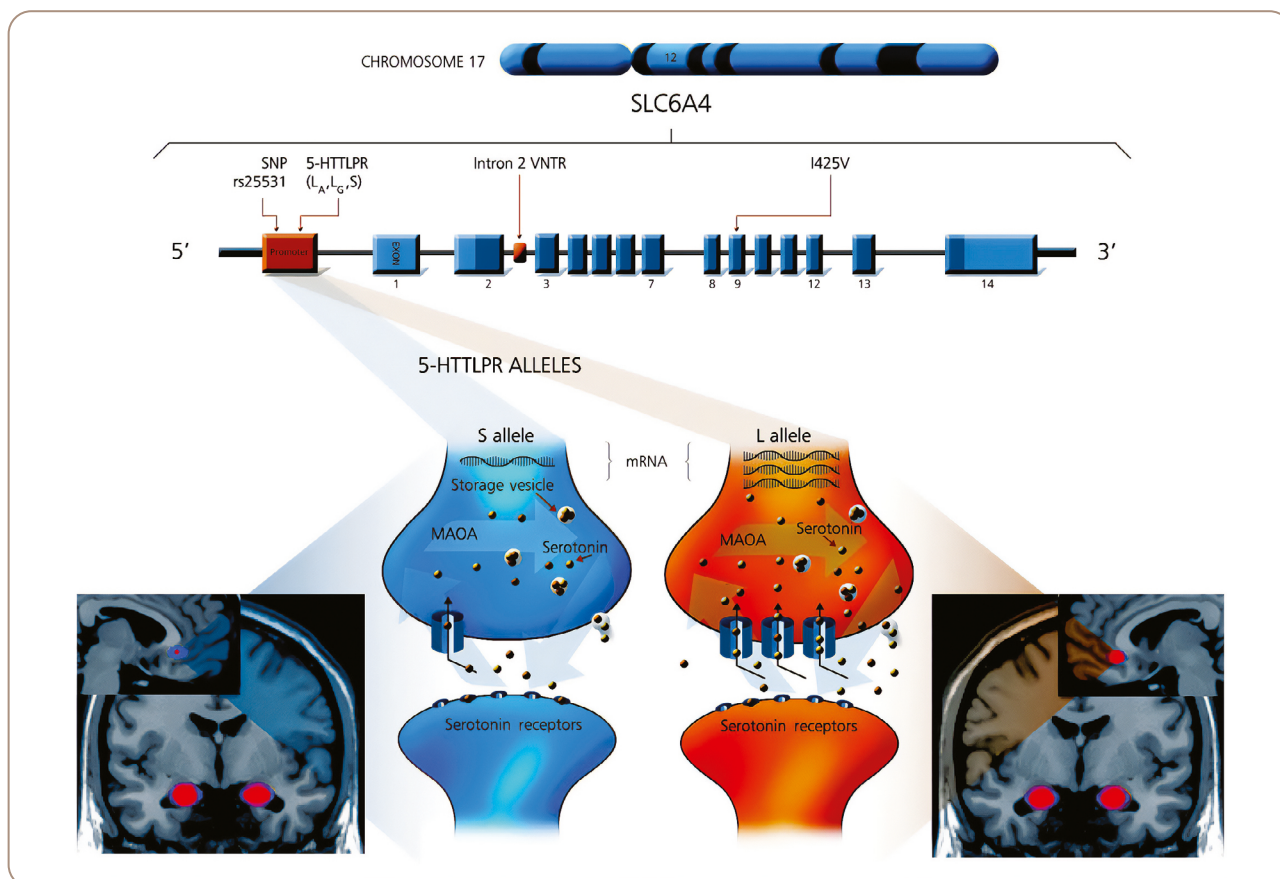


Figure 2. Mapped illustration of the 5-HTTLPR variant in the SLC6A4 gene with allele variants.

Note: 5-HTTLPR — serotonin transporter-linked polymorphic region; CHROMOSOME 17 — the 17th human chromosome, which contains the SLC6A4 gene; Intron 2 VNTR — variable number tandem repeat; MAOA — monoamine oxidase A; mRNA — messenger RNA; rs25531 — is the identifier for a single nucleotide polymorphism (SNP) in the SLC6A4 gene, which affects the expression of the transporter; SNP — single nucleotide polymorphism.

Source: Gerretsen et al., 2009 [21].

It has been reported that there is a link between altered DNA methylation of the gene encoding the serotonin transporter *SLC6A4* and mood disorders, anxiety, as well as amygdala responsiveness [31]. Furthermore, some studies have evaluated the epigenetic changes in the *SLC6A4* gene in schizophrenia patients [32–34]. CpG sites (DNA regions consisting of cytosine and guanine separated by a phosphate) of the *SLC6A4* gene are known to exhibit changes in methylation levels in patients with bipolar disorder [35]. Male patients with schizophrenia also demonstrated similar results [36].

The impact of the 5-HTTLPR variant of the SLC6A4 gene in the development of addictions

Scientific studies in psychogenetics over the past decade have demonstrated that a significant number of mental disorders have a genetic origin [37]. It should be noted

that alcohol abuse is the leading cause of disability and mortality amongst people [38]. The lack of awareness about the harmful effects of alcohol and commitment in society to the ritual of merrymaking, where alcohol is a key element in bringing young people together, can lead to the emergence of behavioral patterns of alcohol consumption [39].

There are two types of addictive disorders:

- chemical addictions (alcohol abuse, drug addiction, toxicomania, etc.);
- non-chemical addictions (pathological gambling, computer addiction, Internet addiction, etc.).

In combination, they can lead to organic disruptions in the higher nervous functions, which can ultimately result in the development of mental disorders [40, 41]. Data suggest that there may be differential genetic vulnerability to alcohol abuse and opiate dependence in serotonergic genes [42].

There is also evidence that the serotonin system plays a role in the pathogenesis of multiple neuropsychiatric disorders and may be involved in addictions such as smoking, since nicotine increases serotonin production in the brain [43–45]. It is assumed that nicotine and other components of tobacco smoke may contain serotonin and thereby contribute to the development of homeostatic resistance [46]. According to some researchers, the genetic variants of different nations lead to different patterns. For example, among residents of Texas (USA) with the LL genotype, smoking was more common than among carriers of the S-allele [47], whereas the 5-HTTLPR variant of the serotonin transporter gene and any association with smoking have not been documented among the Polish population [48].

It is well known that the genetic basis of alcohol abuse lies in the mechanism of ethanol metabolism and the reward system (the neurobiological system associated with dopamine production and the development of addiction) [49]. The scientific community has shown a greater interest in the association between changes in the promoter region of the serotonin transporter gene *SLC6A4* and alcoholism [50]. The S-allele is associated with alcohol consumption, while the L-allele is associated with a positive pharmacological response during the resolution of the withdrawal syndrome [51, 52].

The effects of the 5-HTTLPR variant of the *SLC6A4* gene on the outcomes of therapy with antidepressants in various ethnic groups

SSRIs (citalopram, escitalopram, fluoxetine, fluvoxamine) and serotonin modulators with SSRI-like properties are the main pharmacological options for treating major depressive and anxiety disorders [53–55]. The updated guidelines of the Clinical Pharmacogenetics Implementation Consortium (CPIC) emphasize the import of genotyping *CYP* (*CYP2D6*, *CYP2C19*, *CYP2B6*) genes for dosage optimization; however, our knowledge on the pharmacodynamic *SLC6A4* gene remains insufficient for clinical application [56, 57]. Antidepressants constitute the main therapeutic option for patients with depression; however, about 50% of patients fail to achieve an adequate response to them [58]. The site of action of SSRIs is the serotonin transporter, which means that the concentration of this protein can affect its efficacy both directly and through adaptive changes in the serotonergic function [59, 60]. Due to the differences in the transcriptional activity of 5-HTTLPR,

the dose of SSRIs may inhibit a greater proportion of serotonin in individuals carrying the S-allele, leading to a rapid accumulation of synaptic serotonin and increasing the risk of adverse reactions [20]. The Biallelic (5-HTTLPR) and triallelic (5-HTTLPR/rs25531) patterns in the *SLC6A4* gene are frequently studied, but any idea of association with the antidepressant response remains tenuous [61]. Researchers note differences in the response to SSRIs depending on ethnic variations in 5-HTTLPR: the S-allele is associated with a better antidepressant response in Koreans and Japanese, while the L-allele is associated with a better response in Europeans. However, it is unclear whether the 5-HTTLPR variant and its high expression variant rs25531 have any association with the response to antidepressants [62].

DISCUSSION

Brief interpretation of the results

The 5-HTTLPR variant of the *SLC6A4* gene may interact with the environment and affect the development of addictive disorders [59, 60]. Such stressful events as losses or unfavorable household conditions may have a greater effect on patients with the S-allele, making them predisposed to addictive behavior [63]. It has also been shown that the presence of the S-allele may lead to a reduction in serotonin concentration in synapses, which, in turn, is associated with an increased predisposition to the development of mental disorders and addictive behavior [63, 64]. At a physiological level, this may manifest as emotional instability and increased sensitivity to stress [15, 65]. This emphasizes the importance of considering both genetic and environmental factors when assessing the risk of developing dependences [61, 66]. As can be seen in Table 1, the distribution of genotypes (LL, SL, SS) and alleles (L/S) of the 5-HTTLPR variant of the *SLC6A4* gene varies greatly across different ethnic groups. For instance, in Asian populations (Japanese, Chinese, Yakut), the S-allele predominates (70.6%–80.9%), whereas in European populations (Russian, Ukrainian, Belarusian), the L-allele is more frequently encountered (38.5%–66.7%) [26]. These differences indicate the need to consider population specificity when analyzing genetic risks [26]. Fundamental studies have not identified any unique neurobiological markers (e.g., features of neuroimaging or immune parameters) that would clearly distinguish carriers of the S-allele from patients with other genetic profiles [20].

Table 1. The frequencies of the genotypes and alleles of the 5-HTTLPR variant in the SLC6A4 gene in various populations [26]

Population	n	Frequency of genotypes, % (n)			Frequency of alleles (%)		Reference
		LL	SL	SS	L	S	
Russian (St. Petersburg)	908	38.10 (346)	46.69 (424)	15.19 (138)	61.5	38.5	[67]
Ukrainian	60	21.21 (14)	37.87 (25)	40.90 (27)	61.5	38.5	
Belarusian	39	46.15 (18)	41.02 (16)	12.82 (5)	66.7	33.3	
Chuvash	372	24.46 (91)	51.61 (192)	23.92 (89)	50.3	49.7	
Kabardian	289	26.64 (77)	44.63 (129)	28.71 (83)	49.0	51.0	
Tatar	142	26.05 (37)	51.40 (73)	22.53 (32)	51.8	48.2	
Yakut	158	5.7 (9)	32.3 (51)	62.0 (98)	21.8	78.2	[26]
Chinese (Beijing)	558	6.09 (34)	36.02 (201)	57.88 (323)	24.1	75.9	[44]
Thai	187	9.09 (17)	36.89 (69)	54.01 (101)	27.5	72.5	[20]
Taiwanese	192	10.93 (21)	36.97 (71)	52.08 (100)	29.4	70.6	[68]
Japanese	101	3.7 (4)	31.4 (31)	65.7 (66)	19.3	80.7	[69]
Japanese (Tottori)	501	3.19 (16)	31.73 (159)	65.06 (326)	19.1	80.9	[70]
Chines (Shanghai)	587	6.30 (37)	41.39 (243)	52.29 (307)	27.0	73.0	[71]

Note: The specified samples (Russian — St. Petersburg, Chinese — Beijing, Japanese — Tottori, Chinese — Shanghai) are consistent with the data of original studies (see the references in the table) and reflect local rather than general national samples.

Table 2. Studies of 5-HTTLPR variations of the SLC6A4 gene

Category	Brief description	References
Human studies		
Mental disorders	Relation to schizophrenia, depression, and anxiety in various populations.	[6], [14], [27], [20], [32], [33], [35], [37]
Smoking/nicotine	Association with nicotine dependence and behavioral patterns.	[10], [43–47]
Alcohol	The role of 5-HTTLPR in the development of alcohol abuse.	[19], [51]
Anxiety/stress	Association with panic attacks and stress reactivity.	[17], [21–23], [61], [69], [70]
Pharmacokinetics	Effects on the efficacy of antidepressants (SSRIs).	[53], [63], [67]
Personality/neurodegeneration	Role in personality traits and neurodegenerative processes.	[28], [29], [71]
Population differences	Ethnic variability of alleles and risks.	[25], [42]
Epigenetics	Promotor hypermethylation and its clinical correlates.	[30], [35], [36]
Animal studies		
Epigenetics/environment	The effect of environmental enrichment on SLC6A4 expression and demethylation in mice.	[34]

Discussion of the results

Recently, in the study by Bousman et al., the authors excluded the *SLC6A4* gene from clinical recommendations due to conflicting data and insufficient evidence for its clinical implementation [56]. However, in their systematic review and meta-analysis, Stein et al. showed that the pathological variant of 5-HTTLPR can serve as a marker for antidepressant treatment outcomes in patients with

mental disorders and may be particularly relevant for the use of SSRIs in individuals of European descent [68]. Laje et al. [69] and Rahikainen et al. [70] demonstrated that male patients with a low-functioning genotype SS 5-HTTLPR/rs25531, who were on SSRIs (citalopram), were at increased risk of violent suicide (bringing to suicide). At the same time, studies conducted in Korean patients with severe depression by Jang et al. showed that carriers of the

SS 5-HTTLPR genotypes had significantly better treatment outcomes, while the genotype containing the G (AG+GG) rs25531 variant was associated with remission only [71]. Despite the fact that this pathological allele is involved in the development of addictive disorders, it cannot serve as a clinical marker due to a lack of evidence. Moreover, at the current stage of research, many investigators associate this genetic variant with other mental disorders, such as depression and anxiety (Table 2) [72, 73].

Limitations

Although the coverage of scientific publications based on the keywords used in MEDLINE and eLIBRARY.RU could be considered comprehensive, the descriptive nature of some publications prevented us from including them in the study. The limitation of the search by the specified search engines and keywords led to the heterogeneity of the study material in the meta-analyses, as well as to the retrospective nature of the meta-analyses themselves and the insufficient comprehensiveness of the studies initially selected for them. In this review, only one gene *SLC6A4* and its two variants (5-HTTLPR and rs25531 A<G) were considered. Furthermore, since the pleiotropic effects of the *SLC6A4* gene are associated with depression and anxiety, this limits the possibility of isolated interpretation of its role in the development of addictive disorders. The authors acknowledge the limitations of the information presented and recognize that, even with the most thorough possible approach, the study cannot encompass all aspects of the topic being considered.

CONCLUSION

This review attempted to systematize the data on the role of the 5-HTTLPR variant of the *SLC6A4* gene in the development of addictive disorders, highlighting its ambiguous nature and pleiotropic effects. In contrast to previous studies, the emphasis here is centered on the need for a multidimensional approach to risk assessment that takes into account genetic, environmental, and ethnic factors. Further studies with an in-depth analysis of the molecular mechanisms of the interaction between the 5-HTTLPR variant of the *SLC6A4* gene and the serotonergic system are needed. Future research should also include the development of personalized prevention and treatment strategies, which can potentially improve the efficacy of addiction treatment and reduce the frequency of adverse reactions.

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Information about the authors

***Alexey Vasilievich Krylov**, Junior Researcher, Laboratory of Hereditary Disorders, Department of Molecular Genetics, Yakut Science Centre of Complex Medical Problems; eLibrary SPIN-code: 5746-3015; ORCID: 0009-0005-5977-5518 E-mail: alexkrulovwork@gmail.com

Nadezhda Ivanovna Pavlova, MD, Cand. Sci. (Biology), Leading Researcher, Head of Laboratory of Hereditary Disorders, Department of Molecular Genetics, Yakut Science Centre of Complex Medical Problems; eLibrary SPIN-code: 6167-5254; ORCID: 0000-0001-7862-1876

Alexey Alexeevich Bochurov, Junior Researcher, Laboratory of Hereditary Disorders, Department of Molecular Genetics, Yakut Science Centre of Complex Medical Problems; eLibrary SPIN-code: 1853-0018; ORCID: 0009-0008-5414-4102

*corresponding author

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