NON-COMMUTING OPERATORS AND SPECTRAL INVARIANCE: NON-HERMITIAN COMPLEX HAMILTONIAN AND NON-HERMITIAN REAL HAMILTONIAN

Biswanath Rath

Department of Physics, North Orissa University, Baripada, Orissa, INDIA E.mail:biswanathrath10@gmail.com

ABSTRACT

We notice spectral invariance exists between non-commuting, non-Hermitian (PT-

symmetry) Hamiltonian and corresponding real non-Hermitian (T-symmetry) operators. As an example we consider an analytically solvable model using the unified algebraic meth of (H.B. Zhang, G.Y. Jiang and G.C. Wang J. Math. Phys 56,072103 (2015)) and numerical models. Interestingly the T-symmetry operators are generated using $x \rightleftharpoons p$.

1. **INTRODUCTION**

Since the developments of Quantum Mechanics [1], spectral analysis plays a major role in understanding beauty behind quantum formulations. One of the fundamental quantum postulates is that" commuting operators reflect wave function invariance" mathematically

$$[H_1, H_2] = 0 \tag{1}$$

Here $H_1 \rightarrow E_1$; $H_2 \rightarrow E_2$ and $E_1 \neq E_2$. However till now no such literature which

deals with spectral invariance with

$$[\mathrm{H}_1,\mathrm{H}_2] \neq 0 \tag{2}$$

having different nature of wave functions. However in this paper we would like to report that above relation can yield spectral invariance nature, when recasted as

$$[H_1, (\mathbf{x}, \mathbf{p}), H_2 = H_1(\mathbf{x} \rightleftharpoons \mathbf{p})] \neq 0$$
(3)

In order to address the above relation we discuss real spectra in complex systems [2]. As such it is an abstract mathematics. However subsequent mathematical findings using Quantum Mechanics are verified experimentally both in real analysis [1] and so also in complex analysis [2]. In fact complex nature of quantum mechanics are basically due to space-time (PT) invariant nature of Hamiltonians satisfying the condition [3]

$$PTH (PT)^{-1} = H$$
(4)

Here the complex operator H is not necessarily be hermitian i.e.

$$H \neq H^{\dagger}$$
 (5)

In above P stands for parity operator or space reflection operator having the behaviour

$$PpP^{-1} = -p \tag{6}$$

$$PxP^{-1} = -x \tag{7}$$

Similarly T stands for time - reversal operator

$$TxT^{-1} = -x \tag{8}$$

$$TpT^{-1} = -p \tag{9}$$

where x (co-ordinate) and p(momentum) satisfies the commutation relation

$$[\mathbf{x}, \mathbf{p}] = \mathbf{I} \tag{10}$$

Similarly T-symmetry operator means

$$THT^{-1} = H \tag{11}$$

Mathematically T-symmetry operator can be written as

$$[H, T] = 0$$
 (12)

In this paper we present few model operators on PT-symmetry in nature and generate corresponding T symmetry operator and study the spectral nature as follows.

II. ORIGIN OF T-SYMMETRY OPERATOR

The interesting part of the above symmetry is that on replacing

$$\mathbf{x} \to \mathbf{p}; \mathbf{p} \to \mathbf{x} \tag{13}$$

the magnitude of commutation relation does not change i.e

$$[p, x] = -i \to |[p, x]| = 1 \tag{14}$$

This property has been highlighted in the case of Hermitian operators [4]. However in the case of complex operator

$$ix \rightarrow ip$$
 (15)

the left hand side is PT symmetry and complex in nature, however the right hand side becomes real. Physically a complex PT-symmetry operator is equivalent to real nonhermitian operator.

III. ANALYTICALLY SOLVABLE MODEL

Let us consider the model operator discussed earlier by Bender and Boettecher [2] as

$$H_1 = p^2 + x^2 + ix$$
(16)

having energy eigenvalues

$$E_n = 2n + \frac{5}{4}$$
 (17)

On using $x \rightleftharpoons$, the above operator can be written as

$$h_1 = p^2 + x^2 + ip$$
(18)

In order to find the energy eigenvalues, we use the general relation as follows. For general quadratic operator [5]

$$H_{Ref} = h_{11}p^2 + h_{22}x^2 + ih_{12}(xp + px) + ih_1p + h_2x$$
(19)

having energy eigenvalue

$$\in_{n} = (2n+1)\sqrt{(h_{11}h_{22} + h_{12}^{2})} + \frac{h_{1}^{2}h_{22} - h_{2}^{2}h_{11} - 2h_{1}h_{2}h_{12}}{4(h_{11}h_{22} + h_{12}^{2})}$$
(20)

Following the above we have

 $h_{11} = H_{22} = 1; h_{12} = 0; h_1 = 2, h_2 = 0$ (21)

Under the aboce constraints

$$\in = 2n + \frac{5}{4} \tag{22}$$

It is easy to see that

$$\in = 2n + \frac{5}{4} = E_n \tag{23}$$

Hence we conclude that under the change $x \rightleftharpoons p$ the spectra remains invariant.

IV. NON-ANALYTICAL MODELS

In this case we consider few non-analytical model PT symmetry operators and corresponding T symmetry operators. In all the cases we use matrix diagonalization method to present spectral nature [4, 6] In the present, we solve the eigenvalue relation

$$H|\Phi\rangle = E|\Phi\rangle \tag{24}$$

where

$$|\Phi\rangle = \sum_{m} A_{m} |m\rangle$$
⁽²⁵⁾

and |m> is the harmonic oscillator wave function, satisfying the relation

$$< m|H_0|m> = < m[p^2 + x^2]|m> = (2m + 1)$$
 (26)

A-1 : Broken spectra

The Hamiltonian considered are the following

 $H_2 = p^2 + |x| + ix$ (27)

$$h_2 = x^2 + |p| + ip \tag{28}$$

A-2: Unbroken spectra

The Hamiltonian considered are the following

$$H_3 = p^2 + x^4 + ix$$
(29)

$$h_3 = x^2 + p^4 + ip$$
 (30)

A-3: Broken spectra

The Hamiltonian considered are the following

 $H_4 = p^2 + x^4 + i10x \tag{31}$

$$h_4 = x^2 + p^4 + i10p \tag{32}$$

A-5: Broken spectra

The Hamiltonian considered are the following

$$H_5 = p^2 + ix |x|$$
 (33)

$$h_5 = x^2 + ip |p|$$
 (34)

Few eigenvalues are tabulated in Table-1

V. CONCLUSION

In this paper a reader will notice that all PT-symmetry operator is associated with T-symmetry operator reflecting iso-spectral nature. Similarly our previous study on Hermitian operator also reflects the same thing [5] i.e. every Hermitian operator is associated with another Hermitian operator having the same spectra. In conclusion non-commuting operators generated using $x \Rightarrow p$ will always yield same spectral nature (see figs 1-4). As stated above the above Hamiltonians correspond to different nature of wave functions with a view to justify this nature, we consider the unbroken spectra and display the wave function mod square in fig-5 and 6.

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Table-1: PT-symmetry and T-symmetry operator

n	H ₁	h1	See Fig
0	1.25	1.25	
1	3.25	3.25	
2	5.25	5.25	
3	7.27	7.25	
4	9.25	9.25	
n	H ₂	h2	See Fig
0	1.488158	1.488158	Fig-1
1	3.852053 + 0.868850 i	3.852053 + 0.868850 i	
2	3.852053 – 0.868850 i	3.852053 – 0.868850 i	
3	5.637605 + 1.914109 i	5.637605 + 1.914109 i	
4	5.637605 – 1.914109 i	5.637605 – 1.914109 i	
n	H3	h3	See Fig
0	1.194489	1.194489	Fig-2
1	3.813357	3.813357	
2	7.476329	7.476329	
3	11.661074	11.661074	
4	16.275800	16.275800	
n	H4	h4	See Fig
0	7.919338 – 7.067323 i	7.919338 – 7.067323 i	Fig-3
1	7.919338 + 7.067323 i	7.919338 + 7.067323 i	
2	13.386047 – 3.187097 i	13.386047 – 3.181097 i	
3	13.386047 + 3.181097 i	13.386047 + 3.181097 i	
4	17.957084	17.957084	
n	H5	h5	See Fig
0	1.285090	1.258090	Fig-4
1	4.991283 – 0.780527 i	4.991283 – 0.780527 i	
2	4.991283 + 0.780527 i	4.991283 + 0.780527 i	
3	8.618015 + 3.363276 i	8.618015 + 3.363276 i	
4	13.386047 + 3.181097 i	13.386047 – 3.181097 i	
5	17.957084	17.957084	



Real, $H_2 = p^2 + |x| + ix$ and $h_2 = x^2 + |p| + ip$





Real, $H_3 = p^2 + x^4 + ix$ and $h_3 = x^2 + p^4 + ip$



Figure 2 : Energy eigenvalues



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Figure 5 : Wave function mod square

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Figure 6 : Wave function mod square