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# Factorization solution of a family of quantum nonlinear oscillators 

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#### Abstract

In a recent paper, Cariñena J F, Perelomov A M, Rañada M F and Santander M (2008 J. Phys. A: Math. Gen. 41 085301) analyzed a non-polynomial one-dimensional quantum potential representing an oscillator which they argued was intermediate between the harmonic and isotonic oscillators. In particular they proved that it is Schrödinger soluble, and explicitly obtained the wavefunctions and energies of the bound states. In this paper we show that these results can be obtained much more simply by noting that this potential is a supersymmetric partner potential of the harmonic oscillator. We then use this observation to generate an infinite set of potentials which can exactly be solved in a similar manner.


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## 1. Introduction

In a recent paper in this journal, Cariñena, Perelomov, Rañada and Santander [1], which we refer to from now on as CPRS, investigated the solutions to the eigenvalue problem for the Schrödinger equation ${ }^{1}$

$$
-\frac{\mathrm{d}^{2} \psi}{\mathrm{~d} x^{2}}+\left[x^{2}+8 \frac{2 x^{2}-1}{\left(2 x^{2}+1\right)^{2}}\right] \psi=E \psi
$$

Using some rather involved mathematics they were able to show that the eigenfunctions are given by

$$
\psi_{n}(x)=\frac{P_{n}(x)}{\left(2 x^{2}+1\right)} \mathrm{e}^{-x^{2} / 2}, \quad \text { where } \quad n=0,3,4,5 \ldots,
$$

[^0]and the polynomial factors $P_{n}(x)$ are related to the Hermite polynomials by
\[

P_{n}(x)= $$
\begin{cases}1 & n=0 \\ H_{n}(x)+4 n H_{n-2}(x)+4 n(n-3) H_{n-4}(x) & n=3,4,5 \ldots\end{cases}
$$
\]

The corresponding eigenvalues are given by

$$
E_{n}=-3+2 n \quad \text { where } \quad n=0,3,4,5 \ldots
$$

In this paper we show that the CPRS potential is a supersymmetric partner potential of the harmonic oscillator. This allows us to rederive all the above results in a much simpler fashion. In addition, by considering the question of which other potentials are supersymmetric partners of the harmonic oscillator, we construct an infinite set of exactly soluble potentials, along with their eigenfunctions and eigenvalues.

The rest of the paper is organized as follows. In section 2 we give a brief summary of the ideas of the factorization approach to the Schrödinger equation and supersymmetric partner potentials. In section 3 we prove that the CPRS potential is a partner potential of the harmonic oscillator, and use this to rederive the formulae for its eigenfunctions and eigenvalues. In section 4 we show how to find other partner potentials of the harmonic oscillator, and derive an infinite set of exactly soluble potentials. In section 5 we systematically derive the eigenvalues and eigenfunctions of these new generalized CPRS potentials, whilst in section 6 we make our summary and conclusions.

## 2. The factorization approach

In this section we provide a self-contained introduction to the factorization approach, and the idea of supersymmetric partner potentials. More details of the general theory can be found in [2] and [3]. A discussion of the supersymmetric connection between harmonic and isotonic oscillators can be found in [4].

We start with the one-dimensional single-particle Schrödinger equation,

$$
H_{1} \psi(x)=\left[-\frac{\mathrm{d}^{2}}{\mathrm{~d} x^{2}}+V_{1}(x)\right] \psi(x)=E \psi(x)
$$

The idea is to factorize the Hamiltonian operator, $H_{1}$, which is a second-order differential operator, into a product of two first-order differential operators,

$$
H_{1}=A^{\dagger} A
$$

where

$$
A=\frac{\mathrm{d}}{\mathrm{~d} x}+W(x), \quad A^{\dagger}=-\frac{\mathrm{d}}{\mathrm{~d} x}+W(x)
$$

Upon direct substitution we see that

$$
V_{1}(x)=W^{2}(x)-W^{\prime}(x)
$$

We now define the operator $H_{2}=A A^{\dagger}$ by reversing the order of $A$ and $A^{\dagger}$. Simple algebra shows that $H_{2}$ is a Hamiltonian corresponding to a new potential $V_{2}(x)$,

$$
H_{2}=A A^{\dagger}=-\frac{\mathrm{d}^{2}}{\mathrm{~d} x^{2}}+V_{2}(x), \quad V_{2}(x)=W(x)^{2}+W^{\prime}(x)
$$

The potentials $V_{1}(x)$ and $V_{2}(x)$ are known as supersymmetric partner potentials.
The key result we will be using in later sections is that the eigenvalues and eigenfunctions of $H_{1}$ and $H_{2}$ are related. Suppose that $\psi_{n}^{(1)}$ is an eigenfunction of $H_{1}$ with energy eigenvalue $E_{n}^{(1)}$. Then $A \psi_{n}^{(1)}$ is an eigenfunction of $H_{2}$ with energy eigenvalue $E_{n}^{(1)}$ since

$$
H_{2}\left[A \psi_{n}^{(1)}\right]=A A^{\dagger} A \psi_{n}^{(1)}=A\left[H_{1} \psi_{n}^{(1)}\right]=A\left[E_{n}^{(1)} \psi_{n}^{(1)}\right]=E_{n}^{(1)}\left[A \psi_{n}^{(1)}\right]
$$

The only way that $A \psi_{n}^{(1)}$ could fail to be an eigenfunction of $H_{2}$ is if it equals zero, in which case $W(x)=-\frac{\mathrm{d}}{\mathrm{d} x} \ln \psi_{n}^{(1)}(x)$, which could be true for at most one value of $n$. Similarly suppose that $\psi_{n}^{(2)}$ is an eigenfunction of $H_{2}$ with eigenvalue $E_{n}^{(2)}$. Then $A^{\dagger} \psi_{n}^{(2)}$ is an eigenfunction of $H_{1}$ with energy eigenvalue $E_{n}^{(2)}$ since

$$
H_{1}\left[A^{\dagger} \psi_{n}^{(2)}\right]=A^{\dagger} A A^{\dagger} \psi_{n}^{(2)}=A^{\dagger}\left[H_{2} \psi_{n}^{(2)}\right]=A^{\dagger}\left[E_{n}^{(2)} \psi_{n}^{(2)}\right]=E_{n}^{(2)}\left[A^{\dagger} \psi_{n}^{(2)}\right]
$$

The only way that $A^{\dagger} \psi_{n}^{(2)}$ could fail to be an eigenfunction of $H_{1}$ is if it equals zero, in which case $W(x)=\frac{\mathrm{d}}{\mathrm{d} x} \ln \psi_{n}^{(2)}(x)$, which again could be true for at most one value of $n$.

It follows that, with the possible exception of one level, $H_{1}$ and $H_{2}$ have the same energy spectra. The eigenfunctions of $H_{2}$ can be found from those of $H_{1}$ by applying the operator $A$; the eigenfunctions of $H_{1}$ can be obtained from those of $H_{2}$ by applying the operator $A^{\dagger}$. If $H_{1}$ has one more energy level than $H_{2}$, this energy eigenfunction will be the solution of $A \psi_{0}^{(1)}=0$, and have energy eigenvalue zero; if $H_{2}$ has one more energy level than $H_{1}$, this energy eigenfunction will be a solution of $A^{\dagger} \psi_{0}^{(2)}=0$, and have energy eigenvalue zero. The upshot of all this is that, if we know how to exactly solve one of $H_{1}$ or $H_{2}$, we can immediately derive the exact solution of the other.

Finally we note that we can easily normalize the newly obtained eigenfunctions. If $\psi_{n}^{(1)}$ is a normalized eigenfunction of $H_{1}$ with energy $E_{n}^{(1)}$, the corresponding normalized eigenfunction of $\mathrm{H}_{2}$ is

$$
\psi_{n}^{(2)}=C_{n} A \psi_{n}^{(1)},
$$

where $C_{n}$ is a constant to be determined. If we take the scalar product of this eigenfunction with itself we obtain

$$
1=\left(\psi_{n}^{(2)}, \psi_{n}^{(2)}\right)=\left|C_{n}\right|^{2}\left(\psi_{n}^{(1)}, A^{\dagger} A \psi_{n}^{(1)}\right)=\left|C_{n}\right|^{2} E_{n}^{(1)}
$$

This leads to the result

$$
\psi_{n}^{(2)}=\left[E_{n}^{(1)}\right]^{-1 / 2} A \psi_{n}^{(1)}=\left[E_{n}^{(1)}\right]^{-1 / 2}\left[\frac{\mathrm{~d}}{\mathrm{~d} x}+W(x)\right] \psi_{n}^{(1)},
$$

and we similarly find that

$$
\psi_{n}^{(1)}=\left[E_{n}^{(2)}\right]^{-1 / 2} A^{\dagger} \psi_{n}^{(2)}=\left[E_{n}^{(2)}\right]^{-1 / 2}\left[-\frac{\mathrm{d}}{\mathrm{~d} x}+W(x)\right] \psi_{n}^{(2)}
$$

## 3. Solution of the CPRS potential

We can solve the CPRS potential by noticing that it is a partner potential of the harmonic oscillator. To see this we simply take

$$
W(x)=x+\frac{4 x}{\left(2 x^{2}+1\right)}
$$

from which trivial calculus gives us

$$
\begin{aligned}
& V_{1}(x)=W(x)^{2}-W^{\prime}(x)=x^{2}+8 \frac{\left(2 x^{2}-1\right)}{(2 x+1)^{2}}+3 \\
& V_{2}(x)=W(x)^{2}+W^{\prime}(x)=x^{2}+5
\end{aligned}
$$

If follows that

$$
\begin{array}{ll}
H_{1}=A^{\dagger} A=H_{C}+3 \quad & \rightarrow \quad E_{k}^{(1)}=E_{k}^{C}+3 \\
H_{2}=A A^{\dagger}=H_{H}+5 \quad & \rightarrow \quad E_{k}^{(2)}=E_{k}^{H}+5
\end{array}
$$

where

$$
H_{C}=-\frac{\mathrm{d}^{2}}{\mathrm{~d} x^{2}}+x^{2}+8 \frac{\left(2 x^{2}-1\right)}{\left(2 x^{2}+1\right)^{2}}
$$

is the CPRS Hamiltonian, and

$$
H_{H}=-\frac{\mathrm{d}^{2}}{\mathrm{~d} x^{2}}+x^{2}
$$

is the harmonic oscillator Hamiltonian.
Now the eigenvalues and unnormalized eigenfunctions of the harmonic oscillator [5] are given by

$$
E_{k}^{H}=2 k+1 \quad \text { and } \quad \phi_{k}^{H}(x)=H_{k}(x) \mathrm{e}^{-x^{2} / 2}
$$

where $k=0,1,2 \ldots$ and the Hermite polynomials $H_{k}(x)$ are given by the Rodrigues formula

$$
H_{k}(x)=(-1)^{k} \mathrm{e}^{x^{2}} \frac{\mathrm{~d}^{k}}{\mathrm{~d} x^{k}}\left[\mathrm{e}^{-x^{2}}\right]
$$

It follows trivially that the eigenvalues and unnormalized eigenfunctions of $\mathrm{H}_{2}$ are

$$
E_{k}^{(2)}=2 k+6 \quad \text { and } \quad \psi_{k}^{(2)}(x)=H_{k}(x) \mathrm{e}^{-x^{2} / 2}
$$

The corresponding eigenvalues of $H_{1}$ are therefore $E_{k}^{(1)}=2 k+6$, with unnormalized eigenfunctions

$$
\psi_{k}^{(1)}(x)=A^{\dagger} \psi_{k}^{(2)}(x)=\left[-\frac{\mathrm{d}}{\mathrm{~d} x}+x+\frac{4 x}{(2 x+1)}\right] H_{k}(x) \mathrm{e}^{-x^{2} / 2} .
$$

We can simplify this expression by repeatedly using the Hermite polynomial identities

$$
\begin{aligned}
H_{k}^{\prime}(x) & =2 k H_{k-1}(x) \\
2 x H_{k}(x) & =H_{k+1}(x)+2 k H_{k-1}(x)
\end{aligned}
$$

which can be derived directly from the Rodrigues formula, to obtain

$$
\psi_{k}^{(1)}(x)=\frac{1}{2}\left[H_{k+3}(x)+4(k+3) H_{k+1}(x)+4 k(k+3) H_{k-1}(x)\right] \frac{\mathrm{e}^{-x^{2} / 2}}{\left(2 x^{2}+1\right)} .
$$

The normalization factor for the harmonic oscillator eigenfunctions is

$$
N_{k}^{(2)}=\left[\frac{1}{2^{k} k!\sqrt{\pi}}\right]^{1 / 2},
$$

so the corresponding factor for the CPRS eigenfunctions is

$$
N_{k}^{(1)}=\left[E_{k}^{(2)}\right]^{-1 / 2} N_{k}^{(2)}=\left[\frac{1}{2^{k+1} k!(k+3) \sqrt{\pi}}\right]^{1 / 2}
$$

We have therefore exactly solved the CPRS Hamiltonian, and the eigenvalues and normalized eigenfunctions are given by
$E_{k}^{C}=2 k+3$
$\phi_{k}^{C}(x)=\left[\frac{(k+1)(k+2)}{2^{k+3}(k+3)!\sqrt{\pi}}\right]^{1 / 2} \frac{\left[H_{k+3}(x)+4(k+3) H_{k+1}(x)+4 k(k+3) H_{k-1}(x)\right]}{\left(2 x^{2}+1\right)} \mathrm{e}^{-x^{2} / 2}$
where $k=0,1,2 \ldots$ To compare with the CPRS results we set $n=k+3$ so that $E_{n}^{C}=2 n-3$
$\phi_{n}^{C}(x)=\left[\frac{(n-1)(n-2)}{2^{n} n!\sqrt{\pi}}\right]^{1 / 2} \frac{\left[H_{n}(x)+4 n H_{n-2}(x)+4 n(n-3) H_{n-4}(x)\right]}{\left(2 x^{2}+1\right)} \mathrm{e}^{-x^{2} / 2}$,
where $n=3,4,5 \ldots$ Note that there is one final state possible, which is the solution of $A \phi_{0}=0$, and this will have energy -3 if it exists. Solving this equation gives

$$
\psi_{0}^{C}(x)=C \frac{\mathrm{e}^{-x^{2} / 2}}{\left(2 x^{2}+1\right)}
$$

which is of exactly the same form as the previous $\phi_{n}^{C}(x)$ with $n=0$. Even the form of the normalization constant suggested by placing $n=0$ in the previous equation turns out to be correct, as we show in appendix A. It follows that the final solution of the CPRS Hamiltonian is
$E_{n}^{C}=2 n-3$
$\phi_{n}^{C}(x)=\left[\frac{(n-1)(n-2)}{2^{n} n!\sqrt{\pi}}\right]^{1 / 2} \frac{\left[H_{n}(x)+4 n H_{n-2}(x)+4 n(n-3) H_{n-4}(x)\right]}{\left(2 x^{2}+1\right)} \mathrm{e}^{-x^{2} / 2}$,
where $n=0,3,4,5 \ldots$
We have therefore reproduced the results of CPRS in a very direct and systematic manner by using the fact that the CPRS potential is a partner potential of the harmonic oscillator potential.

## 4. Partner potentials of the harmonic oscillator

The solution of the CPRS Hamiltonian in section 3 was motivated by the observation that it is a partner of the harmonic oscillator. Let us now turn this around, and ask which Hamiltonians, $H_{1}$, are partners to the harmonic oscillator, $V_{2}(x)=x^{2}$, and can thus be solved by the method of section 3. The $W(x)$ needed in the factorization method would then have to satisfy

$$
\frac{\mathrm{d} W}{\mathrm{~d} x}+W(x)^{2}=V_{2}(x)-\lambda
$$

where we have included an irrelevant constant $\lambda$. This is an example of a Riccati differential equation [6], first analyzed in 1724. The method of solution is to substitute

$$
W(x)=\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi(x)=\frac{1}{\phi(x)} \frac{\mathrm{d} \phi}{\mathrm{~d} x},
$$

which leads to the equation

$$
-\frac{\mathrm{d}^{2} \phi}{\mathrm{~d} x^{2}}+V_{2}(x) \phi(x)=\lambda \phi(x)
$$

We see that $\phi(x)$ is a solution of the original Schrödinger equation, in our case the harmonic oscillator. We seem to have gone round in a complete circle, which is not surprising since the central idea of the partner potential method is a mapping between second-order linear and first-order nonlinear differential equations.

We also seem to run into a problem in that every solution $\phi_{k}(x)$ of the original Schrödinger equation, other than the ground state, will have at least one zero $x_{0}$. The $W(x)$ generated from this will then have $\mathrm{a}\left(x-x_{0}\right)^{-1}$ singularity, and then $V_{1}(x)$ will have $\mathrm{a}\left(x-x_{0}\right)^{-2}$ singularity. This approach would therefore seem to only generate soluble potentials which have a singularity at finite $x$, which might be considered an unphysical feature.

This problem can be overcome in the special case of the harmonic oscillator, for which the Schrödinger equation is

$$
-\frac{\mathrm{d}^{2} \phi}{\mathrm{~d} x^{2}}+x^{2} \phi=\lambda \phi
$$

If we set $x=\mathrm{i} y$, this becomes

$$
-\frac{\mathrm{d}^{2} \phi}{\mathrm{~d} y^{2}}+y^{2} \phi=-\lambda \phi
$$

which is the original equation with the irrelevant change in constant $\lambda \rightarrow-\lambda$. In other words, if we set $x \rightarrow \mathrm{i} x$ in the harmonic oscillator eigenfunctions, we get perfectly good $\phi(x)$ which can then generate $W(x)$ and finally $H_{1}$ and $H_{2}$. These $\phi(x)$ would not be good eigenfunctions as they are not normalizable-they behave like $\mathrm{e}^{x^{2} / 2}$ at large $x$-but this is not relevant here. The solutions $\phi(x)$ are thus

$$
\phi_{p}(x)=\mathcal{H}_{p}(x) \mathrm{e}^{x^{2} / 2}
$$

where the pseudo-Hermite polynomials are given by

$$
\mathcal{H}_{p}(x)=(-i)^{p} H_{p}(i x)=\mathrm{e}^{-x^{2}} \frac{\mathrm{~d}^{p}}{\mathrm{~d} x^{p}}\left[\mathrm{e}^{x^{2}}\right] .
$$

They are basically the Hermite polynomials, where the signs of all coefficients are made positive. The even solutions $\phi_{2 m}(x)$ have no real zeros; the odd solutions $\phi_{2 m+1}(x)$ have their only real zero at $x=0$ since they are odd. This leads to a $1 / x^{2}$ singularity at $x=0$, which can be regarded as a centrifugal barrier ${ }^{2}$, as in the case of the isotonic oscillator.

Let us examine the first few partner potentials generated in this manner. For $k=0$,

$$
\begin{aligned}
& \phi_{0}(x)=\mathrm{e}^{x^{2} / 2} \\
& W(x)=\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi_{0}(x)=x \\
& V_{1}(x)=x^{2}-1 \\
& V_{2}(x)=x^{2}+1,
\end{aligned}
$$

and this generates the standard ladder operator solution of the harmonic oscillator, since this shows that the harmonic oscillator is a partner to itself shifted by two units of energy. For $k=1$,

$$
\begin{aligned}
& \phi_{1}(x)=x \mathrm{e}^{x^{2} / 2} \\
& W(x)=\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi_{1}(x)=x+\frac{1}{x} \\
& V_{1}(x)=x^{2}+\frac{2}{x^{2}}+1 \\
& V_{2}(x)=x^{2}+3,
\end{aligned}
$$

and the partner potential is an example of the isotonic oscillator. The eigenfunctions of the isotonic oscillator generated by this method,

$$
\psi_{m}^{(1)}(x)=\left[-\frac{\mathrm{d}}{\mathrm{~d} x}+x+\frac{1}{x}\right] H_{m}(x) \mathrm{e}^{-x^{2} / 2}
$$

are only non-singular for odd $m$, and hence the energy spectrum is $E_{k}=2 k+2$ where $k=1,3,5 \ldots$, so that the level spacing is double that of the harmonic oscillator.

For $k=2$,

$$
\begin{aligned}
& \phi_{2}(x)=\left(4 x^{2}+2\right) \mathrm{e}^{x^{2} / 2} \\
& W(x)=\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi_{2}(x)=x+\frac{4 x}{\left(2 x^{2}+1\right)}
\end{aligned}
$$

[^1]\[

$$
\begin{aligned}
& V_{1}(x)=x^{2}+8 \frac{\left(2 x^{2}-1\right)}{\left(2 x^{2}+1\right)^{2}}+3 \\
& V_{2}(x)=x^{2}+5
\end{aligned}
$$
\]

and this is the CPRS potential we have just solved. This is the first new soluble potential generated by this method, since the solutions of the harmonic and isotonic oscillator are well known.

For $k=3$,

$$
\begin{aligned}
& \phi_{3}(x)=\left(8 x^{3}+12 x\right) \mathrm{e}^{x^{2} / 2} \\
& W(x)=\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi_{3}(x)=x+\frac{1}{x}+\frac{4 x}{\left(2 x^{2}+3\right)} \\
& V_{1}(x)=x^{2}+\frac{2}{x^{2}}+8 \frac{\left(2 x^{2}-3\right)}{\left(2 x^{2}+3\right)^{2}}+5 \\
& V_{2}(x)=x^{2}+7
\end{aligned}
$$

which is a variant of the CPRS potential which includes a centrifugal barrier term.
For $k=4$,

$$
\begin{aligned}
& \phi_{4}(x)=\left(16 x^{4}+48 x^{2}+12\right) \mathrm{e}^{x^{2} / 2} \\
& W(x)=\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi_{4}(x)=x+\frac{\left(16 x^{3}+24 x\right)}{\left(4 x^{4}+12 x^{2}+3\right)} \\
& V_{1}(x)=x^{2}+16 \frac{\left(8 x^{6}+12 x^{4}+18 x^{2}-9\right)}{\left(4 x^{4}+12 x^{2}+3\right)^{2}}+7 \\
& V_{2}(x)=x^{2}+9,
\end{aligned}
$$

which is the harmonic potential plus a rational potential which is regular at the origin, and falls off at infinity.

The common feature of these potentials is that they consist of a harmonic term plus an additional rational function which falls off at infinity like a constant times $1 / x^{2}$. For even $p$ this additional term is symmetric and finite at the origin, and leads to a potential which looks like a harmonic well with an attractive dimple. For odd $p$ the additional term is similar to that for even $p-1$, plus a centrifugal barrier term. The latter means that we consider these potentials in the interval $(0, \infty)$, as in the case of the isotonic oscillator.

## 5. Exact solution of generalized CPRS potentials

In this section we will write the exact closed form solutions of the generalized CPRS potentials, $V_{C}^{p}(x)$, which we define to be the partner potentials of the harmonic oscillator generated by the functions $\phi_{p}(x)$ introduced in the last section. The original CPRS potential corresponds to the case $p=2$.

We first deduce an expression for $V_{C}^{p}(x)$. Since

$$
\begin{aligned}
W(x) & =\frac{\mathrm{d}}{\mathrm{~d} x} \ln \phi_{p}(x)=\frac{\phi_{p}^{\prime}(x)}{\phi_{p}(x)}=x+\frac{\mathcal{H}_{p}^{\prime}(x)}{\mathcal{H}_{p}(x)} \\
W^{\prime}(x) & =\frac{\phi_{p}^{\prime \prime}(x)}{\phi_{p}(x)}-\left[\frac{\phi_{p}^{\prime}(x)}{\phi_{p}(x)}\right]^{2}=1+\frac{\mathcal{H}_{p}^{\prime \prime}(x)}{\mathcal{H}_{p}(x)}-\left[\frac{\mathcal{H}_{p}^{\prime}(x)}{\mathcal{H}_{p}(x)}\right]^{2},
\end{aligned}
$$

it follows that
$V_{2}(x)=W^{2}(x)+W^{\prime}(x)=\frac{\phi_{p}^{\prime \prime}(x)}{\phi_{p}(x)}=x^{2}+2 p+1$
$V_{1}(x)=W^{2}(x)-W^{\prime}(x)=V_{2}(x)-2 W^{\prime}(x)=x^{2}+2 \frac{\mathcal{H}_{p}^{\prime}(x)^{2}-\mathcal{H}_{p}(x) \mathcal{H}_{p}^{\prime \prime}(x)}{\mathcal{H}_{p}(x)^{2}}+2 p-1$.
We therefore deduce the formula for the generalized CPRS potential,

$$
V_{C}^{p}(x)=x^{2}+2 \frac{\mathcal{H}_{p}^{\prime}(x)^{2}-\mathcal{H}_{p}(x) \mathcal{H}_{p}^{\prime \prime}(x)}{\mathcal{H}_{p}(x)^{2}}
$$

Since $\mathcal{H}_{p}(x)$ is a polynomial of order $p$, we see that the second term in $V_{C}^{p}(x)$ is a rational function with numerator a polynomial of degree $2 p-2$, and denominator a polynomial of degree $2 p$. As $x \rightarrow \infty$,

$$
V_{C}^{p}(x) \sim x^{2}+\frac{2 p}{x^{2}}
$$

Starting from the original harmonic oscillator eigenvalues, $E_{k}^{H}=2 k+1$, then $E_{k}^{(1)}=$ $E_{k}^{(2)}=2(k+p+1)$ and $E_{k}^{C}=2 k+3$. The unnormalized eigenfunctions are

$$
\begin{aligned}
\psi_{k}^{(1)}(x) & =A^{\dagger} \psi^{(2)}(x)=\left[-\frac{\mathrm{d}}{\mathrm{~d} x}+x+\frac{\mathcal{H}_{p}^{\prime}(x)}{\mathcal{H}_{p}(x)}\right] H_{k}(x) \mathrm{e}^{-x^{2} / 2} \\
& =\left[2 x H_{k}(x)-H_{k}^{\prime}(x)+\frac{\mathcal{H}_{p}^{\prime}(x)}{\mathcal{H}_{p}(x)} H_{k}(x)\right] \mathrm{e}^{-x^{2} / 2} \\
& =\left[H_{k+1}(x)+\frac{\mathcal{H}_{p}^{\prime}(x)}{\mathcal{H}_{p}(x)} H_{k}(x)\right] \mathrm{e}^{-x^{2} / 2} \\
& =\left[\mathcal{H}_{p}(x) H_{k+1}(x)+\mathcal{H}_{p}^{\prime}(x) H_{k}(x)\right] \frac{\mathrm{e}^{-x^{2} / 2}}{\mathcal{H}_{p}(x)} \\
& =\left[\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!}\{k+p+1\} \frac{(k+p-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x)\right] \frac{\mathrm{e}^{-x^{2} / 2}}{\mathcal{H}_{p}(x)}
\end{aligned}
$$

where the details of the final step in the above derivation are given in appendix B. The normalization factor is given by

$$
N_{k}^{(1)}=\left[E_{k}^{(2)}\right]^{-1 / 2} N_{k}^{(2)}=\left[\frac{1}{2^{k+1} k!(k+p+1) \sqrt{\pi}}\right]^{1 / 2}
$$

If we define $n=k+p+1$ we see that the eigenfunctions are given by

$$
\phi_{n}^{C}(x)=N_{n} \frac{P_{n}(x)}{\mathcal{H}_{p}(x)} \mathrm{e}^{-x^{2} / 2}, \quad \text { where } \quad n=p+1, p+2, p+3 \ldots,
$$

the polynomial factors $P_{n}(x)$ are related to the Hermite polynomials by

$$
P_{n}(x)=\sum_{i=0}^{p} \frac{p!}{i!(p-i)!} n \frac{(n-i-1)!}{(n-2 i)!} H_{n-2 i}(x),
$$

and the normalization constant $N_{n}$ is

$$
N_{n}=\left[\frac{(n-1)(n-2) \ldots(n-p)}{2^{n-p} n!\sqrt{\pi}}\right]^{1 / 2}
$$

The corresponding eigenvalues are given by

$$
E_{n}^{C}=-2 p+1+2 n \quad \text { where } \quad n=p+1, p+2, p+3 \ldots
$$

There is one final state possible, which is given by $A \psi_{0}^{(1)}=0$, and has energy $E_{0}^{(1)}=0$, so that $E_{0}^{C}=-2 p+1$. This state is then given by

$$
\phi_{0}^{C}(x)=\frac{1}{\phi_{p}(x)}=N_{0} \frac{\mathrm{e}^{-x^{2} / 2}}{\mathcal{H}_{p}(x)} .
$$

When $p=2 m$ is even, the normalization constant can be deduced from appendix A to be

$$
N_{0}=\left[\frac{(2 m)!2^{m}}{\sqrt{\pi}}\right]^{1 / 2}
$$

which is exactly what we would obtain if we naively set $n=0$ in our equation for the normalization constant $N_{n}$. It follows that for even $p$, all the previous formulae are still correct, but now we have $n=0, p+1, p+2, p+3, \ldots$, so that we have an equidistant level structure but with $p$ levels missing.

In the case where $p$ is odd, every other level starting with the ground state will have a $1 / x$ singularity at $x=0$, and hence will not be square normalizable, and should be eliminated. This gives an equidistant spectrum with twice the spacing of the harmonic oscillator levels, exactly as in the case of the isotonic oscillator. More precisely, since our Hamiltonian has a strongly singular potential in the case where $p$ is odd, we should consider a self-adjoint extension problem. However, since the singularity is of the form $2 / x^{2}$, the restriction of the domain to functions vanishing at the singularity gives rise to an essentially self-adjoint operator with only one self-adjoint extension ${ }^{3}$. This issue has been extensively discussed, for example in the papers of Casahorrán [4] and Lathouwers [7], and in Reed and Simon's book [8]. The result is that we should restrict the Hamiltonian to the interval $(0, \infty)$, and the eigenfunctions should vanish at $x=0$. This is exactly the same situation that occurs in the isotonic oscillator.

## 6. Summary and conclusions

We have obtained the exact solution of the CPRS quantum nonlinear oscillator in an economical fashion using the methods of supersymmetric quantum mechanics to show that it is a partner of the harmonic oscillator. We have then used this approach to define a countably infinite set of generalized CPRS potentials $V_{p}^{C}(x)$, where $p$ is a positive integer, which we then exactly solve in a similar fashion. The case where $p$ is even is the most interesting, since all the generated eigenfunctions are normalizable. When $p$ is odd, half of the generated eigenfunctions must be removed as they are not normalizable.

The key technical observation is that the harmonic oscillator eigenvalue equation is unchanged under the transformation $x \rightarrow \mathrm{i} x$, so the harmonic oscillator eigenfunctions with $x$ replaced by ix can be used to generate the ladder operators $A$ and $A^{\dagger}$ needed to move between supersymmetric partners. More generally one can use any solution of a Schrödinger equation to generate the ladder operators; it does not have to be a good eigenfunction as normalization is not required.

We finally note that the isotonic oscillator eigenvalue equation is also unchanged under the transformation $x \rightarrow \mathrm{i} x$, and this will be the subject of future investigations.

[^2]
## Acknowledgments

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## Appendix A. Normalization of ground-state eigenfunctions

In this appendix we evaluate the normalization integral needed for the ground state eigenfunctions

$$
\begin{equation*}
I_{2 m}=\int_{-\infty}^{\infty} \frac{\mathrm{e}^{-x^{2}}}{\mathcal{H}_{2 m}(x)^{2}} \mathrm{~d} x=\frac{\sqrt{\pi}}{2^{2 m}(2 m)!} \tag{A.1}
\end{equation*}
$$

To evaluate this integral define $J_{p}(x)$ as the indefinite integral

$$
J_{p}(x)=\int \frac{\mathrm{e}^{-x^{2}}}{\mathcal{H}_{p}(x)^{2}} \mathrm{~d} x=\int G_{p}(x)^{2} \mathrm{e}^{-x^{2}} \mathrm{~d} x
$$

where for simplicity we have defined $G_{p}(x)=1 / \mathcal{H}_{p}(x)$. We can derive identities for the $G_{p}(x)$ by taking the identities for $\mathcal{H}_{p}(x)$,

$$
\begin{aligned}
& \mathcal{H}_{p}^{\prime}(x)=\mathcal{H}_{p+1}(x)-2 x \mathcal{H}_{p}(x) \\
& \mathcal{H}_{p}^{\prime}(x)=2 p \mathcal{H}_{p-1}(x)
\end{aligned}
$$

and substituting $\mathcal{H}_{p}(x)=1 / G_{p}(x)$ to obtain

$$
\begin{aligned}
& G_{p}(x)^{2}=2 x G_{p}(x) G_{p+1}(x)-G_{p}^{\prime}(x) G_{p+1}(x) \\
& G_{p}^{\prime}(x) G_{p-1}(x)=-2 p G_{p}(x)^{2} .
\end{aligned}
$$

We can now substitute the first identity into the formula for $J_{p}(x)$ and integrate by parts to obtain

$$
\begin{aligned}
J_{p}(x) & =\int\left(\left[2 x \mathrm{e}^{-x^{2}}\right] G_{p}(x) G_{p+1}(x)-G_{p}^{\prime}(x) G_{p+1}(x) \mathrm{e}^{-x^{2}}\right) \mathrm{d} x \\
& =-G_{p}(x) G_{p+1}(x) \mathrm{e}^{-x^{2}}+\int\left(\left[G_{p}(x) G_{p+1}(x)\right]^{\prime} \mathrm{e}^{-x^{2}}-G_{p}^{\prime}(x) G_{p+1}(x) \mathrm{e}^{-x^{2}}\right) \mathrm{d} x \\
& =-G_{p}(x) G_{p+1}(x) \mathrm{e}^{-x^{2}}+\int G_{p}(x) G_{p+1}^{\prime}(x) \mathrm{e}^{-x^{2}} \mathrm{~d} x \\
& =-G_{p}(x) G_{p+1}(x) \mathrm{e}^{-x^{2}}-2(p+1) \int G_{p+1}(x)^{2} \mathrm{e}^{-x^{2}} \mathrm{~d} x \\
& =-G_{p}(x) G_{p+1}(x) \mathrm{e}^{-x^{2}}-2(p+1) J_{p+1}(x),
\end{aligned}
$$

where in the fourth step we used the second identity with $p$ replaced by $p+1$. Repeating the process and using the formula to evaluate the definite integral gives
$\left.J_{p}(x)\right|_{0} ^{\infty}=\left.\left[-G_{p}(x)+2(p+1) G_{p+2}(x)\right] G_{p+1}(x) \mathrm{e}^{-x^{2}}\right|_{0} ^{\infty}+\left.4(p+1)(p+2) J_{p+2}(x)\right|_{0} ^{\infty}$.
Suppose now that $p=2 m$ is even. The first term on the right-hand side clearly vanishes as $x \rightarrow \infty$, but as $x \rightarrow 0$ the situation is more complicated since $G_{2 m+1}(x)$ has an $O(1 / x)$ singularity at $x=0$. From the identities for $\mathcal{H}_{p}(x)$ we see that $\mathcal{H}_{2 m}(0)=2(2 m-1) \mathcal{H}_{2 m-2}(0)$ from which we may deduce that $\mathcal{H}_{2 m}(0)=(2 m)!/ m$ ! and hence $G_{2 m}(0)=m!/(2 m)!$. The coefficient multiplying $G_{2 m+1}(x)$ as $x \rightarrow 0$ is thus

$$
-\frac{m!}{(2 m)!}+2(2 m+1) \frac{(m+1)!}{(2 m+2)!}=0 .
$$

The $O(1 / x)$ term thus has coefficient zero, and since the next term is $O(x)$, we see that the first term on the right-hand side also vanishes as $x \rightarrow 0$. If we now multiply by 2 to make the region of integration from $-\infty$ to $\infty$, we obtain the result

$$
I_{2 m}=4(2 m+1)(2 m+2) I_{2 m+2}
$$

Since $I_{0}=\sqrt{\pi}$, the result (A.1) follows.

## Appendix B. Proof of two Hermite polynomial identities

In this appendix we prove two identities involving Hermite and pseudo-Hermite polynomials, which we need to give an explicit form for eigenfunctions of generalized CPRS potentials.

The first identity we prove is

$$
\begin{equation*}
\mathcal{H}_{p}(x) H_{k}(x)=\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!} \frac{(k+p-i)!}{(k+p-2 i)!} H_{k+p-2 i}(x), \tag{B.1}
\end{equation*}
$$

of which the first few examples are

$$
\begin{align*}
& 1 \cdot H_{k}(x)=H_{k}(x) \\
& 2 x \cdot H_{k}(x)=H_{k+1}(x)+2 k H_{k-1}(x) \\
& \left(4 x^{2}+2\right) \cdot H_{k}(x)=H_{k+2}+4(k+1) H_{k}(x)+4 k(k-1) H_{k-2}(x) \tag{B.2}
\end{align*}
$$

To prove this identity we will need the identities

$$
\begin{align*}
& 2 x H_{k}(x)=H_{k+1}(x)+2 k H_{k-1}(x) \\
& \mathcal{H}_{p+1}(x)=2 x \mathcal{H}_{p}(x)+2 p \mathcal{H}_{p-1}(x) \tag{B.3}
\end{align*}
$$

which are easily derived directly from the relevant Rodrigues formulae. In fact the identity (B.1) was originally obtained by using the first of (B.3) to derive the examples (B.2), and thence spot the general pattern.

The proof is by induction on $p$. The case $p=0$ is trivially true; the case $p=1$ is simply the first of (B.3). It follows that the formula is true for $p=0$ and $p=1$. Now assume that it is true for all values up to $p \geqslant 2$. The formula for $p+1$ can then be written as

$$
\begin{align*}
\mathcal{H}_{p+1}(x) H_{k}(x) & =2 x \mathcal{H}_{p}(x) H_{k}(x)+2 p \mathcal{H}_{p-1}(x) H_{k}(x) \\
& =\mathcal{H}_{p}(x) H_{k+1}(x)+2 k \mathcal{H}_{p}(x) H_{k-1}(x)+2 p \mathcal{H}_{p-1}(x) H_{k}(x), \tag{B.4}
\end{align*}
$$

using the identities (B.3). The three terms in (B.4) take the form

$$
\begin{aligned}
\mathcal{H}_{p}(x) H_{k+1}(x) & =\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \\
& =\sum_{i=0}^{p+1} 2^{i} \frac{p!}{i!(p+1-i)!}\{p+1-i\} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \\
2 k \mathcal{H}_{p}(x) H_{k-1}(x) & =2 k \sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x) \\
& =\sum_{i=0}^{p} 2^{i+1} \frac{p!}{i!(p-i)!}\{k\} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x)
\end{aligned}
$$

$$
\begin{align*}
2 p \mathcal{H}_{p-1}(x) H_{k}(x) & =2 p \sum_{i=0}^{p-1} 2^{i} \frac{(p-1)!}{i!(p-1-i)!} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x) \\
& =\sum_{i=0}^{p} 2^{i+1} \frac{p!}{i!(p-i)!}\{p-i\} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x) \tag{B.5}
\end{align*}
$$

where we were able to extend the summation limit in the first and last terms because the additional terms are exactly zero in both cases. Combining the last two terms in (B.5) then yields

$$
\begin{align*}
& \sum_{i=0}^{p} 2^{i+1} \frac{p!}{i!(p-i)!}\{k+p-i\} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x) \\
& \quad=\sum_{i=0}^{p} 2^{i+1} \frac{p!}{i!(p-i)!} \frac{(k+p-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x) \\
& \quad=\sum_{i=1}^{p+1} 2^{i} \frac{p!}{(i-1)!(p+1-i)!} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \\
& \quad=\sum_{i=0}^{p+1} 2^{i} \frac{p!}{i!(p+1-i)!}\{i\} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \tag{B.6}
\end{align*}
$$

where in the second step we shifted $i \rightarrow i-1$, and in the third step we were again able to extend the summation limit because the additional term was zero. Adding (B.6) to the first term of (B.5) gives

$$
\begin{aligned}
\mathcal{H}_{p+1}(x) H_{k}(x) & =\sum_{i=0}^{p+1} 2^{i} \frac{p!}{i!(p+1-i)!}\{p+1\} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \\
& =\sum_{i=0}^{p+1} 2^{i} \frac{(p+1)!}{i!(p+1-i)!} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x),
\end{aligned}
$$

which is just (B.1) with $p$ replaced by $p+1$. This completes the proof by induction.
The second identity we prove is

$$
\begin{equation*}
\mathcal{H}_{p}(x) H_{k+1}(x)+\mathcal{H}_{p}^{\prime}(x) H_{k}(x)=\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!}\{k+p+1\} \frac{(k+p-i)!}{(k+p+1-2 i)!} H_{k+p+1-i}(x) \tag{B.7}
\end{equation*}
$$

To prove this we note from (B.1) that

$$
\begin{align*}
\mathcal{H}_{p}(x) H_{k+1}(x) & =\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!} \frac{(k+p+1-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \\
& =\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!}\{k+p+1-i\} \frac{(k+p-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x), \tag{B.8}
\end{align*}
$$

whilst

$$
\begin{aligned}
\mathcal{H}_{p}^{\prime}(x) H_{k}(x) & =2 p \mathcal{H}_{p-1}(x) H_{k}(x) \\
& =2 p \sum_{i=0}^{p-1} 2^{i} \frac{(p-1)!}{i!(p-1-i)!} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x)
\end{aligned}
$$

$$
\begin{align*}
& =\sum_{i=0}^{p-1} 2^{i+1} \frac{p!}{i!(p-1-i)!} \frac{(k+p-1-i)!}{(k+p-1-2 i)!} H_{k+p-1-2 i}(x) \\
& =\sum_{i=1}^{p} 2^{i} \frac{p!}{(i-1)!(p-i)!} \frac{(k+p-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x) \\
& =\sum_{i=0}^{p} 2^{i} \frac{p!}{i!(p-i)!}\{i\} \frac{(k+p-i)!}{(k+p+1-2 i)!} H_{k+p+1-2 i}(x), \tag{B.9}
\end{align*}
$$

where in the fourth step we shifted $i \rightarrow i-1$, and in the final step we were again able to extend the limit of summation. Adding (B.8) and (B.9) together then gives the required identity (B.7).

Note Added in Proof. After this paper was submitted, our attention was drawn to two papers on related topics. Robnik [9] has considered the factorization solutions of the partner potentials of the harmonic oscillator generated by the harmonic oscillator wavefunctions, $\phi_{k}^{H}(x)$, rather than $\phi_{k}^{H}(\mathrm{i} x)$. These obviously have the non-physical feature of $\left(x-x_{0}\right)^{-2}$ singularities at the nodes of the wavefunction $\phi_{k}^{H}(x)$. Ioffe and Nishnianidze [10] developed a general method to construct almost isospectral potentials using SUSY intertwining relations of third order in derivatives. The CPRS potential is one of a class of potentials soluble by their methods.

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[^0]:    1 Note that we define our Hamiltonian to be twice that used by CPRS, so that our energy values are twice theirs. We use this definition so that the coefficient of the second derivative in Schrödinger's equation is -1 , which is the standard choice in the factorization and supersymmetry literature.

[^1]:    ${ }^{2}$ Of course, the term 'centrifugal barrier' does not make physical sense in one dimension. However, such $\ell(\ell+1) / x^{2}$ singularities are often the result of separating a spherically symmetric three-dimensional problem to yield a radial equation. Hence the term 'centrifugal barrier' is often used for such singular terms in a potential.

[^2]:    ${ }^{3}$ We would like to thank one of the referees and a board member of the journal for drawing our attention to this point.

