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Exciton binding energy and effective mass of CsPbCl₃: a magneto-optical

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High magnetic field spectroscopy has been performed on lead chloride-based perovskite, a material which attracts significant interest for photovoltaic and photonic applications within past decades. Optical properties being mainly driven by the exciton states, we have measured the fundamental parameters, such as the exciton binding energy, effective mass and dielectric constant. Among the inorganic halide perovskites, CsPbCl₃ owns the largest exciton binding energy and effective mass. This blue emitting compound has also been compared with lower band gap energy perovskites and other semiconducting phases, showing comparable band gap dependences for binding energy and Bohr radius.

1. INTRODUCTION

Halide perovskite semiconductors have emerged as promising materials for photovoltaic and optoelectronic applications. Within the last decade, the conversion efficiency of the perovskite solar cells has jumped in an impressive manner from 3% to 25% [1-3]. These outstanding results have also fueled interest on these materials in other domains. As a direct band-gap semiconductor, lead halide perovskite exhibits also excellent electronic and emission properties, very interesting for optoelectronic applications as lasers and photodiodes [4-7], photodetectors [8] or polaritonic devices [9-13]. In addition, the flexible synthesis of nanostructures such as nanocrystals or nanoplatelets, offers numerous opportunities for their integration in quantum devices and the development of applications in the field of quantum optics [14,16].

Emission properties being driven by the excitonic state, a direct experimental determination of the exciton parameters such as binding energy, effective mass and Landé factor are of prime importance for current understanding and future technological applications. Indeed, the theoretical models describing the exciton fine structure or describing the exciton magnetic

behaviour are based on the direct knowledge of the exciton parameters [17-22]. From the experimental point of view, getting accurate values of the binding energy implies to perform and analyse spectroscopic studies on these materials based on techniques such as absorption, photoluminescence or reflectivity. Once the excitonic parameters deduced, they allow the interpretation of results obtained in more complex optical experiments such as time-resolved or photoinduced Faraday rotation measurements [23,24]. From the point of view of applications, the binding energy has to be compared with the thermal energy at the device operating point; it will determine whether the exciton will be stable or dissociates into electron and hole free carriers. The exciton stability is also important for polaritonic devices and room temperature single photon sources or lasers, while for photovoltaic devices, the exciton dissociation is more interesting.

Here, we centre our study on CsPbCl₃ bulk material, which shows the highest energy absorption edge in the family of the all-inorganic lead halide perovskites (CsPbX₃ with X = Cl, Br or I). Recently, lasers and polaritonic effects have been evidenced in this material [9,25,26]. Moreover, nanostructured CsPbCl₃ materials extend the emission energy to the ultraviolet range and allow considering the realization of ultraviolet lasers that

hold important applications from high-resolution bio-imaging and laser therapy to optical storage [27]. Other paths are explored like doping CsPbCl₃ quantum dots with paramagnetic transition metal Mn²⁺ ions, and opening new potentialities to the exciton energy transfer and electro-optical and magnetic properties for this material [28-30]

In this study, we implement low-temperature transmission spectroscopy in pulsed magnetic fields up to 68 T giving directly access to the basic exciton parameters, such as the binding energy of the neutral free exciton and the excitonic reduced effective mass, μ , for CsPbCl₃ films. We found a large exciton binding energy and a weak effective mass making CsPbCl₃ suitable for room temperature laser and polaritonic condensation applications.

2. EXPERIMENTAL RESULTS

CsPbCl₃ perovskite films were grown on a muscovite mica substrate using chemical vapour deposition method described in previous works [25,31-33]. The growth mechanism is known as van-der-Waals epitaxy and needs substrates, such as mica, whose surfaces are inert due to the predominant absence of dangling bonds.

A typical optical microscopy image of grown films is shown in Figure 1 a). The film thickness is in the 200 to 300 nm range. We observe strongly elongated parallelepipedal shaped domains with lateral dimensions of several tens of μm .

Figure 1 b) shows the optical transmission of CsPbCl₃ films obtained at low temperature, 2K, and zero magnetic field. Two minima are observed in the range 2.9 to 3.2 eV. These two minima correspond to the hydrogen-like exciton states $n = 1$ and 2 , respectively at 2.992 eV and 3.040 eV. Assuming an hydrogenic model for the exciton, we can write the energy of the n exciton state as a function of the band-gap energy, E_g , and the exciton binding energy, E_X , as follows:

$$E_n = E_g - \frac{E_X}{n^2} \quad (1)$$

where $E_X = \frac{\mu e^4}{2(4\pi\epsilon_0\epsilon_r\hbar)^2} = E_H \frac{(\frac{\mu}{m_0})}{\epsilon_r^2}$ with e the elementary charge, $\mu = (m_e m_h)/(m_e + m_h)$ the exciton effective mass, $m_{e(h)}$ the electron (hole) effective mass, m_0 the free electron mass, $\hbar$ the reduced Planck's constant, ϵ_0 the vacuum permittivity, ϵ_r the relative permittivity experienced by the exciton and $E_H = 13.605 \text{ eV}$ the binding energy for the hydrogen atom. We deduce $E_X = 64 \pm 1.5 \text{ meV}$ from $E_2 - E_1 = 3E_X/4 = 48 \pm 1 \text{ meV}$. This value is in good agreement with those calculated by DFT methods [34] and experimental determinations [35,36]. Finally, from the energy position of the first minima in the transmission spectrum, we obtain the energy gap for CsPbCl₃: $E_g = 3.056 \text{ eV}$. Figure 1 c) shows the optical transmission of CsPbCl₃ films obtained at room temperature (RT) by using a PerkinElmer Lambda 950 UV-VIS-NIR spectrometer. Only one minimum is observed at RT corresponding to the ground-state exciton transition in agreement with the obtained binding energy of 64 meV, which is larger than the thermal energy at RT. The minimum appears at higher energy than the one observed at 2K, 3.035 eV instead of 2.992 eV, and it is also broader.

E_X is given in Table 1 and compared with the experimental values obtained in other organic and inorganic halide perovskite materials by using the same experimental method. A discussion

is included later below concerning data in Table 1 and Figure 3.

Fig. 1. (a) Typical optical microscopy (reflection configuration) image of a CsPbCl₃ ~ 250 nm thick film grown on muscovite mica following the method reported in references [32] and [33]. (b) Optical transmittance of the film shown in (a) at 2 K and zero magnetic field. S₁ and S₂ correspond respectively to the $n = 1$ and 2 exciton states in the hydrogenic model. (c) Optical transmittance of the film shown in (a) at RT.

Table 1 Exciton parameters in the family of halide perovskite compounds deduced from magneto-optical experiments at 2K.

$\text{CH}_3\text{NH}_3^+ = \text{MA}$ (MethylAmmonium) or $\text{CH}(\text{NH}_2)_2^+ = \text{FA}$ (FormAmidinium).

compound	E_g (eV)	E_X (meV)	m (m_0)	ϵ_r	g_{eff}
^a CsPbCl ₃	3.056	64 ± 1.5	0.202 ± 0.010	6.56 ± 0.24	0.8
^b CsPbBr ₃	2.342	33 ± 1	0.126 ± 0.01	7.3	
^b CsPbI ₃	1.723	15 ± 1	0.114 ± 0.01	10.0	
^c FAPbBr ₃	2.233	22	0.115	8.42	
^c FAPbI ₃	1.501	14	0.09	9.35	2.3
^c MAPbBr ₃	2.292	25	0.117	7.5	
^c MAPbI ₃	1.652	16	0.104	9.4	^d 1.2

- a) This work
- b) ref [37]
- c) ref [38]
- d) ref [39]

Transmission measurements obtained at 2 K in high-magnetic fields are shown in Figure 2 a). These measurements were performed in Faraday configuration with pulsed magnetic fields up to 68 T (pulse duration 500 ms) in Toulouse LNCMI. An optical fiber was used to deliver white light from a Xenon lamp. Focusing spot radius was 100-200 μm allowing excitation of several domains. The circular polarization was introduced *in situ*. Rotation between σ^+ and σ^- polarized light was done by changing the magnetic field direction. The transmitted light was reflected back into a collecting fiber and a spectrometer equipped with a nitrogen cooled CCD detector, which was synchronized to the magnetic field pulse.

Fig. 2. (a) Optical transmittance of the CsPbCl₃ film shown in Fig. 1 a), at 2 K for different magnetic field values. The pulsed magnetic field is perpendicular to the film (Faraday configuration). (b) The energy position of S₁ ($n = 1$) and S₂ ($n = 2$) versus magnetic field. Parabolic fits to the data according to equation (2) in the main text lead to $g_{eff} = 0.8$ and diamagnetic shift coefficients σ_X^n of $0.64 \mu\text{eV}/T^2$ and $2.0 \mu\text{eV}/T^2$ for $n=1$ and 2 , respectively.

In a magnetic field, the exciton energy is modified and can be written as:

$$E_{\pm}^n = E_0^n \pm \frac{1}{2} g_{eff} \mu_B B + \sigma_X^n B^2 \quad (2)$$

where E_0^n is the unperturbed n -exciton energy, g_{eff} is the effective exciton Landé factor, μ_B is the Bohr magneton, B is the magnetic field amplitude and σ_X^n is the diamagnetic shift coefficient. The second term in equation (2) is the linear Zeeman splitting induced by the application of a magnetic field that lifts the exciton states degeneracy and the third term represents the diamagnetic shift that grows quadratically with the field amplitude (in the so-called low field limit).

Fitting simultaneously the energy position of $n = 1$ and 2 state by formula (2), we determine the exciton g -factor $g_{eff} = 0.8$ and the diamagnetic shift coefficients $\sigma_X^1 = 0.64 \pm 0.05 \mu\text{eV} \cdot T^{-2}$ and $\sigma_X^2 = 2.0 \pm 0.2 \mu\text{eV} \cdot T^{-2}$. The diamagnetic shifts are significantly lower than those reported for lower bandgap perovskites [22, 37], which is a natural consequence of a higher exciton binding energy and lower Bohr radius for CsPbCl₃. Additionally the g -factor is significantly lower than in case of iodine or bromide based perovskite.

When the cyclotron energy is significantly smaller than the exciton binding energy, σ_X^1 can be written as a function of the effective exciton mass and the dielectric constant, as follows:

$$\sigma_X^1 = \frac{(4\pi\epsilon_0\epsilon_r)^2 \hbar^4}{4\mu^3 e^2} = \sigma_H \frac{\epsilon_r^2}{(\mu/m_0)^3} \quad (3)$$

$\sigma_H = 1,231 \cdot 10^{-4} \mu\text{eV}/T^2$ is the diamagnetic shift for the hydrogen atom. Combining the expressions of E_X and σ_X , we can obtain the exciton reduced mass and the relative permittivity of CsPbCl₃ as follows:

$$\frac{\mu}{m_0} = \sqrt{\frac{\sigma_H E_H}{\sigma_X E_X}} \quad (4)$$

$$\epsilon_r = \left[\frac{\sigma_H E_H^3}{\sigma_X E_X^3} \right]^{1/4} \quad (5)$$

We then obtain $\frac{\mu}{m_0} = 0.202 \pm 0.010$ and $\epsilon_r = 6.56 \pm 0.24$. Note that, using the numerical calculation of Makado *et al.* [40] for the hydrogenic states under magnetic field, it is possible to deduce the parameter

$\gamma = \frac{\hbar\omega_c}{2E_X}$ at high field, from the behavior of the $n = 1$ and 2 lines splitting. $\omega_c = \frac{eB}{\mu}$ is the exciton cyclotron frequency. One gets $\frac{\gamma}{B} = 4.56 \cdot 10^{-3} T^{-1}$. With $E_X = 64 \text{ meV}$, one obtains $\frac{\mu}{m_0} = 0.199$, in very good agreement with the mass deduced from the $n=1$ state diamagnetic shift in the low-field regime.

Fig. 3. (a) Exciton reduced mass, (b) exciton binding energy and (c) effective dielectric constant as a function of the energy gap. Measurements are done at 2 K. Full orange stars correspond to CsPbX₃ with X = I, Br, or to CsPbI₂Br ref [37]. Empty orange stars correspond to CsPbCl₃ (this work). Open green squares represent results for the MA and FA iodides, bromides or mixed halide (green square at 1.596 eV is for I_{3-x}Cl_x) ref. [38]. Open red squares correspond to MAPb_{1-x}Sn_xI₃ ref [41]. Black solid line in (a) is a linear fit to the data. Vertical dashed green line indicates the value of the energy gap at which a maximum of the effective dielectric constant is found for the considered perovskite compounds.

The comparison of the exciton reduced mass, exciton binding energy and effective dielectric screening for perovskites with different halides is presented in Figure 3. The exciton binding energy increases with increasing bandgap as in more conventional semiconductors (see also Figure 4). At the same time, it can be seen that the effective exciton mass follows reasonably the prediction of a simple two band $k \cdot p$ model, $\mu \propto E_g$ [22]. Recent DFT calculations have addressed the electron and hole mass determination in CsPbX₃ compounds and predicted a significant enhancement of the carrier effective mass when iodine or bromine are exchanged by chlorine due to weaker hybridization of halide orbitals with lead [34,42-44]. With the increase of the halide atom mass the energetic position and space extension of s and p orbitals favor the overlap with the Pb²⁺ states increasing the band dispersion and therefore decreasing the effective exciton mass when one moves from the lighter Cl to heavier Br and I atoms. This effect is especially visible in the valence band, which contains significant admixture of halide orbitals. While most of these theoretical studies have underestimated the exciton mass as compared to experimental measurements, an increase in the exciton mass has been predicted from iodide to chloride, with a ratio of the exciton

masses between 1.53 and 2.24. These predicted ratios are in agreement with the experimental value found in this study, and leading to a ratio $0.202 / 0.114 = 1.77$. Moreover, as observed in Table I, organic and inorganic compounds with the same halide atom possess very similar exciton mass. Based on this observation, the comparison of our results with calculations on methyl ammonium lead chloride should be pertinent. Sendner *et al.* [45] have performed DFT calculations taking into consideration exchange-correlation functional and spin-orbit coupling, and have predicted an exciton mass of $0.20 m_0$ for MaPbCl_3 very similar to the one we measured in CsPbCl_3 .

Fig. 4. The exciton binding energy E_x (a) and the Bohr radius a_x (b) versus the gap energy of halide-based perovskites and other semiconductors with different crystalline structures, either Zinc Blende, Wurtzite or Diamond. The solid lines correspond to fits with respectively $E_x \propto E_g^{1.62}$ and $a_x \propto E_g^{-1.33}$. Perovskite values are given in Table 1. InAs and ZnO [46], GaAs [47], ZnTe [48], ZnS [49], AlN [50] and for the other semiconductors [51].

The dielectric screening is shown in Figure 3c. It follows the expected trend, *i.e.*, at low temperature when the motion of cations is frozen the metal halide cage determine in first approximation the screening properties. The dielectric constant then decreases when the bromide or iodine is exchanged with lighter Cl atoms [37,41]. Such a substitution increases the energy of phonon modes [43] since this energy is proportional to the inverse of reduced mass of atoms participating in vibration. As we can see in Figure 3c, the substitution of Pb atom by Sn atom results also in a decrease of the dielectric screening. Then, Figure 3c shows that the effective dielectric constant decreases as the reduced mass of atoms building the crystal decreases and consequently a non-monotonous behavior of the dielectric constant *versus* the energy gap of the considered perovskite compounds.

Putting together arguments given above and the latter discussion, we underline that the substitution of I^- anions by Cl^- increases the exciton binding energy by two ways, by increasing

carriers effective mass and by reducing dielectric screening. We also see in Figure 3 b) that when substituting Pb^{2+} cation by Sn^{2+} the exciton binding energy does not vary significantly. That is the result of a compensation of two effects: (i) the reduction of the effective mass that participates to the decrease of the exciton binding energy and, (ii) the reduction of the dielectric screening that leads to the increase of the exciton binding energy.

To finish, we have compared the exciton binding energy E_x and the Bohr radius a_x of the halide-based perovskites with other semiconductor families, either those showing Zinc blende, Wurtzite or Diamond crystal structures. As shown in Figure 4, the perovskite parameters follow the trends observed for other semiconducting crystals and phases, on a large range of the band gap energy, with a large increase of E_x and decrease of a_x *versus* E_g . The solid lines correspond to least-square fits with $E_x \propto E_g^{1.62}$ and $a_x \propto E_g^{-1.33}$. As related previously, in a simple two-band model, $\mu \propto E_g$. Chadi and Cohen [52] have shown, on a large band gap energy, that the static dielectric constant follows a law $\epsilon_r - 1 \propto a_0^{-1.5} E_g^{-1}$, with the lattice parameter a_0 . At the same time, Dalven [53] has evidenced an empirical law $E_g \propto a_0^{-2}$ on a large band gap range. All these laws lead to the relations: $E_x \propto E_g^{1.5}$ and $a_x \propto E_g^{-1.25}$, with exponents very close to the ones obtained in Figure 4.

3. CONCLUSION

We have performed a detailed magneto-optical study of the all-inorganic halide perovskite CsPbCl_3 . The analysis of the transmission spectra obtained at low temperature and pulse magnetic field up to 68 T allows the determination of the crucial parameters to describe the exciton properties without assumption on the strength of the dielectric screening. This experimental approach is then self-consistent. We underline that the values of the exciton binding energy and effective mass increase with the band gap energy for the all-organic or inorganic perovskite compounds meanwhile the permittivity values have a non monotonous behaviour. The large exciton binding energy measured in CsPbCl_3 compounds makes them very promising materials for opto-electronic and polaritonic applications at room temperature.

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Disclosures

The authors declare no conflicts of interest.

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