

Performance of CVD diamond detectors for single ion beam-tagging applications in hadrontherapy monitoring

Sébastien Curtoni, Marie-Laure Gallin-Martel, Latifa Abbassi, Alexandre Bes, Germain Bosson, Johann Collot, Thierry Crozes, Denis Dauvergne, Wout de Nolf, Pierre Everaere, et al.

▶ To cite this version:

Sébastien Curtoni, Marie-Laure Gallin-Martel, Latifa Abbassi, Alexandre Bes, Germain Bosson, et al.. Performance of CVD diamond detectors for single ion beam-tagging applications in hadrontherapy monitoring. Nuclear Instruments and Methods in Physics Research Section A: Accelerators, Spectrometers, Detectors and Associated Equipment, 2021, 1015, pp.165757. 10.1016/j.nima.2021.165757 . hal-03227464

HAL Id: hal-03227464 https://hal.science/hal-03227464

Submitted on 16 Oct 2023 $\,$

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

Version of Record: https://www.sciencedirect.com/science/article/pii/S0168900221007427 Manuscript_9d71408c7f9fb4dc21b08d9b8de93d51

Performance of CVD diamond detectors for single ion beam-tagging applications in hadrontherapy monitoring

S. Curtoni^{a,*}, M.-L. Gallin-Martel^a, S. Marcatili^a, L. Abbassi^b, A. Bes^a, G. Bosson^a, J. Collot^a, T. Crozes^b, D. Dauvergne^a, W. De Nolf^c, P. Everaere^a, L. Gallin-Martel^a, A. Ghimouz^a, F. Haddad^{d,e}, C. Hoarau^a, J.-Y. Hostachy^a, C. Koumeir^{d,e}, A. Lacoste^a, V. Métivier^e, J. Morse^c, J.-F. Motte^b, J.-F. Muraz^a, F. Poirier^{d,e}, F. E. Rarbi^a, O. Rossetto^a, M. Salomé^c, N. Servagent^e, E. Testa^f, M. Yamouni^a
^aUniversité Grenoble-Alpes, CNRS, Grenoble INP, LPSC-IN2P3 UMR 5821, 38000 Grenoble, France
^bUniversité Grenoble-Alpes, CNRS, Institut Néel, NANOFAB UPR2940, 38000 Grenoble, France
^cEuropean Synchrotron Radiation Facility, 38000 Grenoble, France
^dGIP ARRONAX, 44800 Saint Herblain, France

^eUniversité de Nantes, CNRS, IMT Atlantique, SUBATECH-IN2P3 UMR 6457, 44000 Nantes, France ^fUniversité de Lyon, CNRS, IP2I-IN2P3 UMR 5822, 69000 Lyon, France

Abstract

In the context of online ion range verification in particle therapy, the CLaRyS collaboration is developing Prompt-Gamma (PG) detection systems. The originality in the CLaRyS approach is to use a beam-tagging hodoscope in coincidence with the gamma detectors to provide both temporal and spatial information of the incoming ions. The ion range sensitivity of such PG detection systems could be improved by detecting single ions with a 100 ps (σ) time resolution, through a quality assurance procedure at low beam intensity at the beginning of the treatment session. This work presents

Preprint submitted to NIM A

July 18, 2021

^{*}Corresponding author: curtoni@cppm.in2p3.fr

the investigations that led to assessment of the Chemical Vapor Deposition (CVD) diamond detectors performance to fulfil these requirements. A 90 Sr beta source, 68 MeV protons, 95 MeV/u carbon ions and a synchrotron X-ray pulsed beam were used to measure the time resolution, single ion detection efficiency and proton counting capability of various CVD diamond samples. An offline technique, based on double-sided readout with fast current preamplifiers used to improve the signal-to-noise ratio, is also presented. The different tests highlighted Time-Of-Flight resolutions ranging from $13 \text{ ps}(\sigma)$ to $250 \,\mathrm{ps}$ (σ), depending on the diamond crystal quality and the particle type and energy. The single 68 MeV proton detection efficiency of various large area polycrystalline (pCVD) samples was measured to be >96% using coincidence measurements with a single-crystal reference detector. Single-crystal CVD (sCVD) diamond proved to be able to count a discrete number of simultaneous protons while it was not achievable with a polycrystalline sample. Considering the results of the present study, two diamond hodoscope demonstrators are under development: one based on sCVD, and one of larger size based on pCVD. They will be used for the purpose of single ion as well as ion bunches detection, either at reduced or clinical beam intensities. *Keywords:* CVD diamond, hadrontherapy, ion range verification, time resolution, detection efficiency, particle counting, beam monitoring

1 1. Introduction

Hadrontherapy is an external radiotherapy modality based on light ion
beams [1, 2]. Even though the ballistic properties of ions and the enhanced
relative biological effectiveness represent essential advantages of particle ther-

apy compared to conventional X-ray radiotherapy, it is still facing limitations 5 due to ion range uncertainties arising at every stage of the treatment pro-6 cedure [3]. They currently lead physicians to set ion range specific safety 7 margins that limit the dose conformation and prevent them to plan irradia-8 tion fields where organs at risk are located close beyond the targeted volume. 9 In this context, several experimental approaches have been developed to 10 build an online ion range verification system [4, 5]. Among them, prompt-11 gamma-based verification techniques [6] propose to retrieve the actual ion 12 range from the emission profile of prompt-gamma photons (PG) that are 13 emitted along the ion path by excited target nuclei or ion fragments right 14 after inelastic collisions between incoming ions and target nuclei. To get rid 15 of the inherent and substantial neutron-induced background also produced 16 during these nuclear interactions, PG detection systems use a Time-Of-Flight 17 (TOF) based gamma-neutron discrimination. It is generally carried out by 18 coincidence measurements between the gamma camera trigger and the ion 19 bunch time of arrival given by the accelerator radio-frequency signal (RF). 20 Provided the body-camera distance is set to a few tens of centimeters, an 21 overall TOF resolution of 1 ns (σ) is sufficient to achieve this purpose. 22

Instead of using the accelerator RF as a START signal, the CLaRyS collaboration proposes to set up a beam-tagging hodoscope upstream from the patient at reduced intensity (~ 1 ion/bunch). It will also provide an ion transverse position that is useful for PG vertices reconstruction with PG imaging systems (PGI). The direct detection of incoming ions also makes the TOF measurement independent of the beam time structure and/or any potential RF phase shift as has been observed [7]. According to this idea, the collaboration has developed a $12.8 \times 12.8 \text{ cm}^2$ scintillating-fiber hodoscope. It has been tested and characterized on proton and carbon ion beams [8] and the results highlighted a $0.7 \text{ ns} (\sigma)$ time resolution and a detection efficiency up to 98%.

Considerable improvements can be achieved in the sensitivity of potential 34 ion range shift determination by improving the TOF resolution down to a few 35 hundred picoseconds. This holds for PGI and prompt gamma timing (PGT) 36 [9, 10, 11] and is thoroughly discussed in [12]. Different detector technologies 37 could enable the development of a beam monitor with a 100 ps (σ) time 38 resolution for single ion detection [13, 14, 15]. The collaboration has chosen 39 to focus on Chemical Vapor Desposition (CVD) diamond technology in order 40 to develop a beam hodoscope upgrade combining an excellent time resolution 41 [16], [15] (and references therein) and high radiation hardness guaranteeing 42 long-term stability in clinical conditions. 43

The current work presents investigations led on diamond detectors, at 44 first, to evaluate polycrystalline (pCVD) single proton detection efficiency. 45 Then, experiments were carried out to assess single crystal (sCVD), pCVD 46 and Diamond On Iridium (DOI) detector ability to perform TOF measure-47 ments with a 100 ps resolution, using 68 MeV single protons in ARRONAX 48 (Saint-Herblain, France), 95 MeV/u carbon ions in GANIL (Caen, France), 49 short pulses of 8.53 keV X-rays at ESRF (Grenoble, France) and minimum 50 ionizing particle (MIP) with a ⁹⁰Sr laboratory source. Finally, the sCVD and 51 pCVD diamond detectors single particle counting capabilities have been eval-52 uated with the 68 MeV proton beam delivered by the ARRONAX cyclotron 53 at low intensity (6 pA \sim 1 proton/bunch).

⁵⁵ 2. Material and methods

56 2.1. Detectors assembly and generic experimental set-up

The detector-grade diamond samples used in the present work are com-57 mercially available and produced by Chemical Vapor Deposition (CVD). The 58 sCVD diamonds were purchased from Element6 [17], pCVD diamonds from 59 Element6, II-VI [18] and Diamond Delaware Knives (DDK) [19], and DOI 60 diamonds from Audiatec [20] and Augsburg University. The tested samples 61 ranged from 300 $\mu{\rm m}$ to 500 $\mu{\rm m}$ in thickness, and from 4.5 \times 4.5 ${\rm mm}^2$ to 20 62 \times 20 mm² in area. In particular, large pCVD are foreseen for the assembly 63 of a large size hodoscope. Diamond samples were assembled as pad detec-64 tors as described in [21, 22]. A thin aluminum disk-shaped metallization 65 was performed either by physical evaporation [23] (50 nm) or by sputtering 66 (100 nm). The diamonds were sandwiched between two 50Ω -adapted printed 67 circuit boards (PCB), allowing bias of both polarities and signal readout con-68 nections on both sides. 69

For the different tests presented in this work, diamond detectors were 70 systematically tested by pair. Figure 1 illustrates the general configuration 71 used during the tests. Two diamond detectors are exposed to a particle 72 beam. The detector under test is positioned upstream from a smaller size 73 reference detector (a sCVD sample, unless stated otherwise). For each beam 74 test, both detectors were enclosed together in an aluminum shielding box 75 with front and rear apertures covered with 12 µm-thick aluminized Mylar 76 films [21, 22]. The output channels of the detectors were coupled to broad-77 band amplifiers (CIVIDEC C2-HV [24] or Greenstream DBA III and IV-R) 78 and analog signals were digitized using fast digitizers (a 500 MHz, 3.2 GS/s 79

Table 1: Summary of the various diamond samples tested within this work. DE = Single proton detection efficiency, TOF = Time-Of-Flight resolution, C = counting, $\sigma_t = intrinsic$ time resolution.

Discussed	Provider	Size	Metallization		Computed	Trade Janith	Involved in
Diamond		(mm^3)	Diam. (mm)	Thick. (nm)	capacitance (pF)	Tested with	measurements:
sCVD	E6	$4.5 \times 4.5 \times 0.517$	3	50	0.7	X, β , p, ¹² C	DE, TOF, C
sCVD	E6	$4.5 \times 4.5 \times 0.517$	3	50	0.7	β	TOF
pCVD	E6	$10 \times 10 \times 0.3$	7	50 - 100	6.5	$X, p, {}^{12}C$	DE, TOF
pCVD	E6	$20 \times 20 \times 0.5$	16	50	20	¹² C	TOF
pCVD	II-VI	$10 \times 10 \times 0.5$	7	100	3.9	р	DE, TOF
pCVD	DDK	$5 \times 5 \times 0.3$	3	100	1.2	р	DE, TOF, C
DOI	Augsburg Univ.	$5 \times 5 \times 0.3$	3	50 - 100	1.2	X, p, ${}^{12}C$	TOF, σ_t
DOI	Audiatec	$5 \times 5 \times 0.3$	3	50	1.2	X	σ_t



Figure 1: Generic experimental set-up used for detection efficiency, timing and counting measurements presented in this work. Specific dedicated additions in the set-up are presented in the corresponding subsections.

WaveCatcher digitizer [25] or a 2 GHz, 20 GS/s LeCroy Digital Storage Oscilloscope (DSO)).

⁸² 2.2. Side-to-side signals summation

During the interaction of a ionizing particle in a diamond detector, the 83 same signal S (absolute value) is induced on both electrodes by the elec-84 tron/hole pairs drift. In practice, each side is read by a single preamplifier 85 generating an output signal of amplitude S^{side_i} (with i=1,2) with a corre-86 sponding noise level $\sigma_n^{side_i}$ resulting in a signal-to-noise ratio $S^{side_i}/\sigma_n^{side_i}$. 87 First, we assume that the intrinsic noise generated by the diamond itself is 88 negligible at 300 K compared to that induced by the wide-band preamplifier. 89 Then, the noise of each preamplifier is assumed to be an independent Gaus-90 sian white noise. Therefore, the resulting noise on the sum-signal σ_n^{sum} can 91 be expressed as follows: 92

$$\sigma_n^{sum} = \sqrt{\left(\sigma_n^{side_1}\right)^2 + \left(\sigma_n^{side_2}\right)^2} = \sigma_n^{side_1} \oplus \sigma_n^{side_2} \,. \tag{1}$$

A sum signal $S^{sum} = S^{side_1} - S^{side_2}$ (the two side signals are of opposite polarity) can be derived as well as a sum-signal-to-noise ratio:

$$S/N_{sum} = \frac{S^{side_1} - S^{side_2}}{\sigma_n^{side_1} \oplus \sigma_n^{side_2}}.$$
 (2)

If one supposes now that the two preamplifiers are strictly identical, Equation
2 becomes:

$$S/N_{sum} = \frac{2S^{side}}{\sqrt{2\sigma_n^2}} = \frac{2S^{side}}{\sigma_n\sqrt{2}} = \sqrt{2} \cdot S/N_{side} \,. \tag{3}$$

⁹⁷ Using the sum signal, the signal-to-noise ratio (SNR) can be increased by a ⁹⁸ factor $\sqrt{2}$. By doubling the amplitude of the signal, the slope in the rising ⁹⁹ edge of the sum-signal can become up to twice as much as the one measured on each electrode signal improving the time resolution of the detector consequently. This technique was used for the detection efficiency and timing
measurements presented in Sections 2.3.1 and 2.3.2. Note that this technique
requires identical preamplifiers (same pulse shape), a strict adjustment of the
pulses risetime and a null delay between both side signals.

¹⁰⁵ 2.3. Experimental tests and data analysis procedures

¹⁰⁶ 2.3.1. Single proton detection efficiency of pCVD detectors

The single proton detection efficiency has been evaluated with 68 MeV 107 protons during a dedicated experiment at ARRONAX IBA C70 isochronous 108 cyclotron, Nantes (with a fixed Radio-Frequency of 30.45 MHz) [26, 27]. In 109 order to restrict the incoming beam to bunches containing at most one single 110 proton, the beam intensity was lowered down to 50 fA. Three different pCVD 111 detectors presented in Table 1 and based on samples coming from different 112 providers were tested one-by-one during the experiment in reproducible con-113 ditions. Each pCVD sample was tested in coincidence with the same sCVD 114 reference detector. The detectors box was set up and aligned between two 115 2.5 cm-thick aluminum collimators with 5 mm gaps. The upstream one was 116 used to constrain the proton incidence to the sensitive surface of the smallest 117 detector. The downstream one reduced the beam halo caused by the scatter-118 ing of protons in the PCBs. Behind the second collimator, a PTW T34058 119 gas ionization chamber (IC) and a 5 mm-thick plastic scintillator coupled to 120 a photomultiplier tube (PMT) were aligned with the beam. The IC was 121 coupled to a PTW Unidos electrometer to measure the beam current while 122 the scintillator was used to get a redundant spectroscopic information of in-123 coming ions that was used for the efficiency measurement. The applied bias 124

voltage was +300 V for the Element6 and DDK pCVD detectors, +500V for 125 the II-VI pCVD detector and +300 V for the sCVD detector, according to 126 the scheme presented in Figure 1. The applied biasing was +400 V for the 127 IC and -800 V for the PMT. The two pCVD output channels were coupled 128 to Greenstream DBA IV-R preamplifiers [28] while CIVIDEC C2-HV pream-129 plifiers were used for the sCVD sample. Analog signals from the diamond 130 detectors and the scintillator were sampled using the WaveCatcher digitizer. 131 To assess the single proton detection efficiency of pCVD samples, mea-132 surements in coincidence with the two reference detectors (the sCVD and 133 the scintillator) were used. First, recorded events that corresponded to a 134 double coincidence between the two reference detectors were identified using 135 a coincidence window of duration $\delta t = 1.25$ ns, as well as low and high volt-136 age thresholds selecting single proton events. Among the $N_{double}(\delta t)$ events 137 that corresponded to these criteria, triple and random coincidences were 138 tested event-by-event on the pCVD samples using two coincidence windows 139 and a voltage threshold scanning. The triple coincidence window was the 140 same as the one applied on the reference detectors. The random coincidence 141 window was delayed by 15 ns, between two consecutive bunches (32.84 ns). 142 Using the voltage threshold sweep, a voltage comparison is performed on 143 the pCVD signal between the threshold level V_{th} and the waveform segments 144 contained within the coincidence windows. For each V_{th} value, we counted 145 $N_{triple}(V_{th}; \delta t)$ triple coincidences between the pCVD and the two reference 146 detectors and $N_{random}(V_{th}; \delta t)$ random coincidences triggered by noise fluctu-147 ations in the pCVD signals. Thus, we can define a true coincidence detection 148 efficiency $\epsilon(V_{th}; \delta t)$ at a given threshold value V_{th} (δt is a fixed parameter) 149

150 as follows:

$$\epsilon \left(V_{th}; \delta t \right) = \frac{N_{triple} \left(V_{th}; \delta t \right)}{N_{double} \left(\delta t \right)} \times \left(1 - \frac{N_{random} \left(V_{th}; \delta t \right)}{N_{double} \left(\delta t \right)} \right) \,. \tag{4}$$

Equation 4 can be understood as the product of the probability to detect a true triple coincidence and the probability not to detect a random coincidence. As δt is fixed, the single proton detection efficiency ϵ_{det} was here defined as the maximum of the obtained $\epsilon(V_{th})$ function.

155 2.3.2. Timing resolution with single ions

The experimental set-up presented in Figure 1 was used in various beam test configurations as listed in Section 1 in order to evaluate the TOF resolution achievable between two diamond detectors of various crystalline qualities. Within the scope of this article, the TOF resolution σ_{TOF} of a pair of independent detectors with respective time resolution σ_{t_1} and σ_{t_2} is defined as:

$$\sigma_{TOF} = \sqrt{\sigma_{t_1}^2 + \sigma_{t_2}^2} \,. \tag{5}$$

Timing measurements were carried out on the digitized waveforms using 162 a normalised threshold algorithm, as defined in [29]. Once the amplitude 163 of a pulse is detected, a constant fraction of this value (between 20% and 164 50%) is computed. The pulse time stamp is finally obtained by means of 165 a linear interpolation between waveform samples, thus emulating an ana-166 log Constant Fraction Discrimination (CFD). Unless stated otherwise, the 167 timing resolution is derived from the statistical dispersion measured on the 168 timing difference between the sum signals of the two detectors involved. Fur-169

thermore, a pair of diamond detectors composed of a $5.0 \times 5.0 \times 0.3 \,\mathrm{mm^3}$ DOI detector produced at Augsburg University and a $4.5 \times 4.5 \times 0.517 \,\mathrm{mm^3}$ sCVD detector produced by Element6 were tested together in the different beam tests presented in this work. They were systematically tested to provide a common reference for comparison purposes and are referred as the sCVD-DOI reference pair, later on in this article.

At ARRONAX, the timing measurements were carried out on the wave-176 form datasets we acquired for the single proton detection efficiency assess-177 ment. We could therefore measure the TOF resolution for the three pCVD-178 sCVD couples, as they are presented in Table 1 and Section 2.3.1. For each 179 V_{th} value, the events subset which fulfilled the triple coincidence criterion and 180 did not trigger random coincidences was selected. On this subset, the pulse 181 discrimination was performed using the normalised threshold algorithm on 182 the pCVD and sCVD sum signals. The distribution of the timing difference 183 between the discriminated pCVD and sCVD signals was then stored in a his-184 togram. Since some distributions demonstrated non-gaussian tails on each 185 side, the root mean square (RMS) value of the histogram was chosen as an 186 estimator of the TOF resolution for all histograms (*i.e.* for each V_{th} value). 187

The timing measurements at GANIL were carried out with single 95 MeV/u carbon ions and the standard bench presented in Figure 1. Yet, one noteworthy difference is that only one CIVIDEC C2-HV preamplifier could be used for each diamond detector during this test, preventing us from using the signal summation technique introduced in Section 2.2. The waveforms were digitized with the 3.2 GS/s WaveCatcher system. Two pairs of detectors were tested. The first one is the sCVD-DOI reference pair and the ¹⁹⁵ second one is composed of two Element6 pCVD detectors, $20 \times 20 \times 0.5 \text{ mm}^3$ ¹⁹⁶ and $10 \times 10 \times 0.3 \text{ mm}^3$ respectively. They were metallized as pad detectors, ¹⁹⁷ with disk-shaped 50nm-thick Al electrodes and respective diameter of 16 mm ¹⁹⁸ and 7 mm.

199 2.3.3. Timing resolution with a pulsed X-ray beam

At ESRF, the combined use of a X-ray micro-beam and various atten-200 uators set up upstream from the detectors enabled us to study the time 201 response of a pair of diamond detectors as a function of the energy depo-202 sition. The test beam took place in ID21 beamline [30] that delivered a 203 8.53 keV X-ray micro-beam while the ESRF synchrotron was running in 4-204 bunch mode. In this configuration, the pulsed beam RF was $f_{RF} = 1.42 \text{ MHz}$ 205 $(T_{RF} = 704 \,\mathrm{ns})$ and the bunch duration was 100 ps. With the maximum elec-206 tron beam current (32 mA) circulating in the synchrotron, the primary X-ray 207 flux was $\phi_{32mA} = 1.79 \cdot 10^9$ photons/s [31], which corresponds to $1.26 \cdot 10^3$ pho-208 tons/bunch. The absorption length of X-rays with an energy $E_X = 8.53 \text{ keV}$ 209 in diamond is $1/\mu_{diam} \sim 790 \ \mu m$ [32]. As a result, the energy deposition is 210 almost uniformly distributed over the thickness of the tested samples (300 -211 500 µm) thus mimicking the passage of single charged particles. 212

The two detectors used here were a $4.5 \times 4.5 \times 0.517 \text{ mm}^3$ Element6 sCVD detector and a $5.0 \times 5.0 \times 0.3 \text{ mm}^3$ Audiatec DOI detector. Both were metallized with aluminium disk electrodes of 3 mm diameter. For each attenuator used (Al and Ti foils with various thicknesses), an acquisition of the signals coming from the two detectors as well as of the RF signal was performed. Each side electrode of the Audiatec sensor was coupled to a CIVIDEC C2-HV preamplifier while only one was used on the sCVD detector. For a given attenuator type and thickness, the energy deposits of a X-ray bunch in the DOI and sCVD detectors (thereafter noted ΔE_{DOI} and ΔE_{sCVD}) are computed using Beer-Lambert law as follows :

$$\Delta E_{DOI} = E_X \cdot \frac{\phi_{32mA}}{f_{RF}} \cdot \exp\left(-\mu_{att} x_{att} - \mu_{PET} x_{PET}\right)$$

$$\cdot \left[1 - \exp\left(-\mu_{diam} d_{DOI}\right)\right], \qquad (6)$$

$$\Delta E_{sCVD} = E_X \cdot \frac{\phi_{32mA}}{f_{RF}}$$

$$\cdot \exp\left(-\mu_{att} x_{att} - \mu_{PET} x_{PET} - \mu_{diam} d_{DOI}\right) \qquad (7)$$

$$\cdot \left[1 - \exp\left(-\mu_{diam} d_{sCVD}\right)\right] ,$$

where μ and x are respectively the attenuation coefficient at 8.53 keV and the thickness of the considered material (att = attenuator, PET = Mylar, diam = diamond) while d is the thickness of the detector. In this set-up, the attenuation (< 0.7%) of the beam in the air path between the detectors was neglected. For each acquisition, the signals have been processed using the normalised threshold algorithm at 50%, as defined in Section 2.3.2.

A previous experiment with the DOI detector from Augsburg University 229 had been carried out with only a few number of attenuators. During this 230 test, the Augsburg DOI sample was coupled to two preamplifiers while the 231 sCVD detector was equipped to only one preamplifier. In this case, we had 232 measured the side-to-side time difference between pulses generated on the 233 two electrodes of the DOI detector. We performed the same measurement 234 with the Audiatec sample and compared the timing performance obtained in 235 both cases. 236

237 2.3.4. Timing resolution with MIP electrons

Using MIP-like electrons allowed us to determine a lower bound of the 238 timing resolution that could be obtained for the detection of single particles. 239 In this case, two Element6 sCVD diamond detectors $(4.5 \times 4.5 \times 0.51 \text{ mm}^3)$ 240 each) were used. They were exposed to a collimated beam of electrons from a 241 ⁹⁰Sr source, with an energy up to 2.28 MeV. The source, the collimators and 242 the detectors were all enclosed in a U-shaped rail ensuring the mechanical 243 alignment of the set-up. An assembly of four scintillating fibres coupled to a 244 common PMT was added downstream from the diamond detectors. It was 245 used as an external trigger to detect electrons in the higher energy part of 246 the β spectrum. Both electrodes of the two diamond detectors were coupled 247 to CIVIDEC C2-HV preamplifiers via 10 cm coaxial cables. The signals 248 produced by the four preamplifiers were digitized using a LeCroy HDO9404 249 DSO (4 GHz, 20 GS/s, 10 bits). The applied bias voltage was -500 V on the 250 two diamond detectors. 251

252 2.3.5. Proton counting

The counting and monitoring capabilities of the diamond samples were 253 also tested at ARRONAX. The DDK pCVD detector and the Element6 254 sCVD detector presented in Table 1 were selected for this test. Only one 255 output channel per detector was used here and the biased electrodes of the 256 pCVD and the sCVD detectors were coupled to one preamplifier. In order to 257 acquire 2 µs-long waveforms (corresponding to 60 RF periods at 30.45 MHz), 258 the sampling rate was lowered down to 2.5 GS/s. A 2.5 cm-thick aluminum 250 collimator with a 1 mm gap was set up in front of the detectors to constrain 260 the beam to a section smaller than the sensitive diameter of the detectors. 261

²⁶² Typical signal waveforms acquired simultaneously on the two detectors are²⁶³ shown in Figure 2.



Figure 2: Compared waveforms acquired simultaneously on the Element6 sCVD detector (top, red) and the DDK pCVD detector (bottom, blue) using a 2.5 GS/s sampling rate.

In order to count the number of protons contained in the bunches, charge measurement was performed by numerical integration of the waveforms on both detectors. First, a baseline correction was achieved by projecting all the waveform samples voltage values in an histogram. Considering that some RF periods do not contain protons at this beam current level ($\sim 5 \text{ pA}$) and that the signal duration is short compared to the RF period, the histogram exhibits a dominant Gaussian noise peak that can be fitted to derive its mean

and standard deviation parameters. They are then defined as estimators of 271 the baseline offset value and the noise level σ of the considered waveform, 272 respectively. After subtraction of the obtained offset, each waveform is sub-273 divided in 60 33ns-long segments (corresponding to the 60 RF periods). For 274 each RF period, the numerical integration is done by summing up the sam-275 ples contained in the corresponding segment. The charge response of both 276 detectors can thus be compared on a bunch-by-bunch basis, as presented in 277 Section 3.3. 278

From the counting statistics, it is possible to derive a mean beam current value. Later on in this paper, we will consider that at a given beam current I_{beam} , the number of protons contained in a bunch is a discrete random variable X according to Poisson law, with a λ parameter such as $\lambda \propto I_{beam}$. The probability P(X = k) of having k protons in a bunch is therefore:

$$P(X=k) = \frac{\lambda^k}{k!} e^{-\lambda} \,. \tag{8}$$

In the case of an ideal beam delivering exactly one proton per bunch ($Q_{bunch} = e$) with a period $T_{beam} = 32.84 \,\mathrm{ns}$ (corresponding to the period of the AR-RONAX cyclotron RF signal), the average beam current I_{ref} is given by:

$$I_{ref} = \frac{Q_{bunch}}{T_{beam}} = \frac{1.602 \cdot 10^{-19}}{32.84 \cdot 10^{-9}} = 4.872 \cdot 10^{-12} \text{A s/s} \,. \tag{9}$$

²⁸⁷ Then the average beam current I_{beam} can be derived as follows:

$$I_{beam} = \lambda I_{ref} \,. \tag{10}$$

This expression will be used later on in this work to estimate the average beam current during the counting experiment. Since λ is the parameter of the Poisson law describing a bunch's proton multiplicity, this analysis carried out on time windows corresponding to 60 consecutive bunches results in a standard deviation $\sigma_{\lambda} = \sqrt{\lambda/60}$.

293 3. Results

²⁹⁴ 3.1. Single proton detection efficiency

The results of the analysis presented in Section 2.3.1 and carried out on 295 the sum signals of the three pCVD samples presented on Table 1 are com-296 bined in Figure 3 (dashed lines). The three pCVD detectors highlight the 297 same behaviour according to the V_{th} value. If V_{th} is close to zero, the proba-298 bility of a random coincidence triggered by noise fluctuations is comparable 299 to the probability to trigger on the true event pulse. As a consequence, ϵ 300 remains low. If V_{th} increases, the noise-triggered random coincidence prob-301 ability decreases and ϵ increases. Beyond an optimal V_{th} value for which 302 ϵ is maximized, the threshold starts rejecting true events resulting in the 303 degradation of the detection efficiency. 304

Following Section 2.3.1, the single 68 MeV proton detection efficiency ϵ_{det} is 305 here defined as $\epsilon_{det} = \max(\epsilon(V_{th}))$. For the three pCVD detectors, ϵ_{det} is 306 obtained at $V_{th} \sim 13 \,\mathrm{mV}$ and reaches 98% for the Element6 sample, while 307 97% is obtained in the case of the II-VI and DDK samples. These results are 308 in good agreement with measurements carried out in similar conditions in a 309 previous study [33] and bring an additional information on random triggering 310 probability. As these results depend on the δt parameter, one should note 311 that they could be improved by reducing the coincidence window, which is 312 in principle possible due to the shortness of the analog pulses. In our case, 313



Figure 3: Single 68 MeV proton detection efficiency (dashed lines) and TOF resolution (solid lines) of three pCVD detectors as a function of the threshold value V_{th} used for the pCVD sum signal discrimination. The coincidence window duration is $\delta t = 1.25$ ns.

the 3.2 GS/s sampling rate was the limiting factor since $\delta t = 1.25$ ns only corresponds to four consecutive waveform samples. Besides, we verified that reducing δt induced an increase of the true coincidence detection efficiency, particularly for low V_{th} values. An additional event selection criterion based on time-over-threshold (TOT) could be used to reject high frequency noisegenerated triggers.

- 320 3.2. Timing performance
- 321 3.2.1. 68 MeV protons

Figure 3 also shows the results of the timing measurements that were carried out on the same acquired datasets. On Figure 3, the measured TOF resolution is plotted as a function of V_{th} for the three pCVD detectors (solid

lines). A similar evolution of the TOF resolution can be observed with the 325 three pCVD samples. It can be noticed that as long as the threshold level 326 remains below the value maximizing the detection efficiency, the measured 327 TOF resolution is rather constant. For higher values, as the threshold rejects 328 low amplitude signals, the SNR of the selected events increases. Since the 329 time resolution of diamond detectors is directly related to the SNR [15, 34], 330 the TOF resolution improves as well. In the case of the Element6 detector, 331 the TOF resolution ranges from 220 ps (RMS) to 162 ps (RMS) and 218 ps 332 (RMS) is obtained at best efficiency. The TOF resolution measured with 333 the II-VI samples ranges from 227 ps (RMS) to 172 ps (RMS) (225 ps at best 334 efficiency). The DDK provides the best results with a TOF resolution ranging 335 from 192 ps (RMS) to 139 ps (RMS) (191 ps at best efficiency). 336

The overall better performance obtained with the DDK detector is re-337 lated to the capacitance of the devices. That plays a crucial role in timing 338 measurements [15, 34]. Using the geometries defined in Table 1 and the 339 relative permittivity of diamond ($\varepsilon_r = 5.7$), the DDK detector's computed 340 capacitance is 1.2 pF compared to 6.5 and 3.9 pF for the Element6 and II-VI 341 detectors respectively. Despite that, the Element6 sample's timing response 342 appears to be slightly better than that of the II-VI sample. It therefore tends 343 to show the superior performance of the Element6 pCVD detector compared 344 to the II-VI one. As a comparison, in a previous beam test, the sCVD-DOI 345 reference pair of diamond detectors had been tested in similar conditions and 346 reached a 94 ps (σ) TOF resolution [11]. 347

348 3.2.2. 95 MeV/u carbon ions

The results of the timing measurements carried out at GANIL are pre-349 sented in Figure 4. Due to the large energy deposition in the detectors (25 350 MeV in DOI and 44 MeV in sCVD according to SRIM simulations [35]), 351 the high SNR enabled us to lower the discrimination fraction down to 20%. 352 Thus, the two detector pairs highlighted excellent results. In each case, the 353 distribution could be fitted to derive the σ_{TOF} value. The measured TOF 354 resolution of the sCVD-DOI pair is $\sigma_{TOF} = 13 \text{ ps.}$ In the case of the pCVD 355 pair, the obtained TOF resolution was 66 ps (σ). The difference between the 356 results obtained with the two pairs can be explained by the quality of the 357



Figure 4: Time difference distributions obtained with two pairs of diamond detectors and single 95 MeV/u carbon ions at GANIL The two pairs were the sCVD-DOI reference pair (green) and two large area pCVD detectors (blue).

involved samples and the large size of the pCVD detectors (with computed
capacitances of 20 pF and 6.5 pF). In any case, the two pairs demonstrated
excellent results nicely fitting with the objectives of the hodoscope.

361 3.2.3. Bunches of 8.53 keV synchrotron radiation X-rays

Figure 5-(Top) represents, for each energy deposition (each attenuator), 362 the TOF resolution measured between the two detectors (red) and between 363 each detector and the beam RF (DOI = blue and sCVD = green) as a function 364 of the deposited energy. The results are fitted with a function $\sigma_{TOF} = C/\Delta E$ 365 with C a parameter to highlight the correlation between the TOF resolution 366 and the deposited energy. Due to the low jitter in the beamline RF signal, 367 the TOF measurements using the RF signal and a single diamond detec-368 tor give better results than TOF measurements made between two diamond 369 detectors. It also provides a common reference allowing us to deduce that 370 the sCVD detector gives a better result than the Audiatec one. The re-371 sults obtained with these two DOI detectors (Audiatec and Augsburg) are 372 compared in Figure 5 (Bottom). As the electronic channels used in both 373 cases were identical (1 CIVIDEC C2-HV per channel), the contribution of 374 the electronic jitter is the same for the two detectors. The better timing 375 response of the Augsburg DOI can therefore be related to the intrinsic better 376 performance of the detector, in comparison with the Audiatec one. While 377 the side-to-side jitter evolution measured on the Audiatec detector fits pretty 378 well with an inverse function, it is not the case of the Augsburg DOI sample. 379 The dashed line fit is drawn to show which correlation would be expected 380 with these measurements but they seem to be less sensitive to the energy de-381 position. A possible explanation may rely on the surface heterogeneity of the 382

Augsburg DOI sample, already highlighted in [31], that could explain that the signal shape will depend on the hit position on the detector. Therefore, the jitter of the Augsburg sample could be dominated by other factors than the energy deposition.



Figure 5: (Top) TOF resolution as a function of the energy deposited by a bunch of 8.53 keV X-rays in a pair of diamond detectors composed of a Element6 sCVD and a Audiatec DOI. (Bottom) Standard deviation of the side-to-side pulses time difference measured on the Audiatec DOI detector (red) and the Augsburg University DOI detector (blue) as a function of the energy deposition in the detectors.

387 3.2.4. Minimum Ionizing Particles (β source)

Prior to the timing measurement itself, a preliminary analysis was performed at LPSC. As the acquisition was triggered by the downstream scintillator, it is shown in Figure 6 that one can assess the existing correlation between the responses of the two diamond detectors. Each detector response corresponds to the integral of the sum signals. The result mainly exhibits two



Figure 6: Correlation between the charge response of the two sCVD detectors used for the MIP timing measurement (the acquisition is triggered by the external downstream scintillator). The first peak centered on (0;0) is the noise peak. The second peak is due to single high energy electrons depositing the same amount of energy in the two detectors. The contour drawn on the distribution is the graphical cut applied on the data to measure the time resolution.

distributions. The first one is centered on zero and the second corresponds to a signal measured simultaneously on both detectors. The statistical predominance of the distribution centered on zero is due to the trigger on the external scintillator which has i) a larger area than diamonds, and may then detect electrons outside the diamond active areas, and ii) a low detection threshold, enabling triggering on background.



Figure 7: Distribution of the time difference between the sum signals of the two Element6 sCVD detectors detecting the same high energy β electrons.

In order to measure the time resolution of the detectors, a graphical selection was performed on the data as illustrated by the black contour drawn in Figure 6. Since the electrons of highest energy are close to MIP, their energy deposition in the two detectors is expected to be almost constant. By selecting the events which exhibit the same charge response in both detectors, we can thus select electrons in the higher energy part of the beta spectrum.

The time difference measured on the sum signals of the selected events is 405 presented in Figure 7. Different estimators can then be used to derive the 406 TOF resolution in this case. An optimistic estimation would consist in using 407 the standard deviation given by a Gaussian fit. Choosing such a parameter 408 neglects the tails present on both sides of the distribution. Under these con-409 ditions, $\sigma_{TOF} = 240 \,\mathrm{ps}$ is obtained, which corresponds to a timing resolution 410 of 170 ps for a single detector. A more objective estimator is the RMS of the 411 distribution. This one takes into account its tails that strongly degrade the 412 TOF resolution. Using this estimator, the TOF resolution is 749 ps (RMS), 413 *i.e.* a timing resolution of $530 \,\mathrm{ps}$ (RMS) for one detector. However, these 414 values were obtained using a 10 ns coincidence window, which is of the same 415 order of the signal duration, and may contain random coincidences. 416

417 3.2.5. Summary of Time-Of-Flight measurements

The TOF resolution we measured at laboratory, at ARRONAX and GANIL are summarized in Table 2 where D1 is the upstream detector and D2 is the downstream one.

The correlation between time resolution and energy deposition (and there-421 fore SNR) can be clearly observed. The measurements are better with the 422 sCVD-DOI reference pair. Using carbon ions at GANIL, the performance 423 of the pCVD pair is excellent. Considering the large energy deposition of 424 carbon ions with energies in the hadron therapy range, developing a pCVD 425 holoscope reaching a time resolution $\leq 100 \text{ ps}(\sigma)$ is achievable in carbon ion 426 therapy. Finally, the measurement with beta electrons allowed us to define 427 a lower limit to these TOF resolutions. 428

Table 2: Summary of the different timing measurements presented. The energy deposition of single ions has been estimated with SRIM simulations. *Result from a previous study [11], given here for completeness purpose.

Diamond D1 VS D2	Manuf.	Size (mm ³)	Computed capacitance (pF)	Particle type	Particle energy (MeV)	Energy deposition per particle/pulse (MeV)	Sum signals used ?	Measured TOF resolution $(ps \sigma)$
sCVD sCVD	E6	$4.5 \times 4.5 \times 0.517$	0.7	⁹⁰ Sr decay electron	$\sim MIP$	~ 0.3	1	240 ± 2
	Augsburg	$5 \times 5 \times 0.3$	1.2	proton	68*	1.0 1.6	1	$94.1 \pm 0.4^{*}$
DOI sCVD	E6	$4.5 \times 4.5 \times 0.517$	0.7	carbon ion	1140	25 44	×	12.7 ± 0.2
	Audiatec E6	$5 \times 5 \times 0.3$ $4.5 \times 4.5 \times 0.517$	1.2 0.7	X-ray pulse (no attenuator)	$8.53\cdot 10^{-3}$	3.4 3.3	DOI only	58.3 ± 0.5
	E6 E6	$10 \times 10 \times 0.3$ $4.5 \times 4.5 \times 0.517$	6.5 0.7			1.0 1.6	1	218 ± 1
pCVD sCVD	II-VI E6	$10 \times 10 \times 0.5$ $4.5 \times 4.5 \times 0.517$	3.9 0.7	proton	68	1.6 1.6	1	225 ± 1
	DDK E6	$5 \times 5 \times 0.3$ $4.5 \times 4.5 \times 0.517$	1.2 0.7			1.0 1.6	1	191 ± 1
pCVD pCVD	E6 E6	$20 \times 20 \times 0.5$ $10 \times 10 \times 0.3$	20 6.5	carbon ion	1140	44 25	×	65.7 ± 1.1

429 3.3. Proton counting

The monitoring and counting capabilities of sCVD and pCVD detectors 430 were evaluated at ARRONAX at a beam intensity around 1 proton/bunch. 431 The results of the bunch-generated ionisation charge Q_{bunch} as measured 432 simultaneously on the sCVD and pCVD detectors are presented in Figure 8 433 (Top). On the one hand, it can be clearly observed that the sCVD detector 434 has an energy resolution which is sufficient to distinguish a discrete number 435 of protons contained in each bunch. The 2D distribution therefore exhibits 6 436 peaks corresponding to bunches whose content ranges from 0 to 5 protons. On 437 the other hand, the pCVD detector's energy resolution is not good enough to 438

count the number of protons in the bunch, leading to an overlap of the charge
distributions corresponding to different numbers of protons. The different
charge distributions could be separated in this case thanks to the correlation
with the sCVD detector.

The sCVD Q_{bunch} distribution which corresponds to the X-projection of 443 the 2D histogram in Figure 8-Top is the convolution of a Poisson distribution 444 of parameter λ with the Gaussian response function of the detector. One 445 can fit the whole distribution with the sum of 6 Gauss functions. From the 446 obtained fit parameters and using the fact that $\lambda = (k+1) \cdot P(k+1)/P(k)$, 447 one can derive the actual λ value. From this analysis, an experimental value 448 of $\lambda = 1.26 \pm 0.02$ is obtained, thus resulting in a mean beam current $I_{beam} =$ 449 $6.16 \pm 0.10 \,\mathrm{pA}$ (using Equations 9 and 10). The error corresponds to the 450 RMS of the λ values obtained using the different k values. Moreover, the 451 I_{beam} uncertainty could be easily reduced by increasing the integration time. 452 Note that the method is only valid if the beam current is constant during 453 the acquisition. 454

Nevertheless, the bunch content separation provided by the sCVD detec-455 tor can be used to assess the linearity of the pCVD detector's mean charge 456 response. Fixed thresholds can be set on the sCVD Q_{bunch} distribution so 457 that the response of the pCVD detector can be conditioned by the response of 458 the sCVD detector. For each peak in the sCVD Q_{bunch} distribution (ranging 459 from 0 to 4 protons), the histogram of the corresponding charge measured 460 on the pCVD is drawn and the obtained mean and RMS values are stored. 461 The correlation between the mean responses of the two detectors for each 462 number of protons can thus be plotted (Figure 8 Bottom). In spite of the 463



Figure 8: (Top) Bunch-generated ionisation charge measured on the Element6 sCVD detector as a function of the charge generated in the DDK detector. (Bottom) Mean charge generated in the sCVD detector as a function of the mean charge generated in the DDK detector, for a discrete number of protons in the bunch. Error bars are given in the figure but are hidden by the marker size. They correspond to the statistical error obtained for each number of protons.

⁴⁶⁴ poor pCVD energy resolution, one can note that its mean charge response ⁴⁶⁵ remains linear with the number of protons contained in the bunch. There ⁴⁶⁶ is no evidence of charge-saturation, and this suggests that pCVD detectors ⁴⁶⁷ could be used at higher beam currents (typically clinical beam currents) to ⁴⁶⁸ provide an efficient beam monitoring, where the proton bunch multiplicity ⁴⁶⁹ prevents from counting the protons individually.

470 4. Discussion

At first, measurements were carried out to evaluate diamond single proton 471 true coincidence detection efficiency, *i.e.* the probability to detect a proton in 472 a time coincidence window as short as possible (1.25 ns), without triggering 473 on the noise, in order to perform efficient TOF measurements on any incident 474 proton. They were done using 68 MeV protons in a single incident particle 475 mode (50 fA). In this way, we could make measurements independent from 476 the beam time structure. Three pCVD diamond sensors were tested. A pro-477 ton coincidence detection efficiency > 96% is reached on the three diamond 478 samples. To perform such a measurement, diamond detectors were read out 479 on both sides which, in the case of an off-line data analysis, makes it possible 480 to increase the SNR by a factor $\sqrt{2}$ when using identical read-out channels. 481 If one is using this method online, particular care should be paid onto the 482 exact synchronization and identical pulse shapes on the two readout chan-483 nels. Indeed, we could observe that if a slight delay between the two signals 484 is not corrected, the time resolution is degraded. Also, if the noise levels 485 are different on the two signals, the noise level of the sum signal is domi-486 nated by the worse level as expected from Equation 1, which degrades the 487

performance obtained with the best readout channel. This has an effect on both efficiency and timing resolution. In the case of a single channel reading, data analysis has shown that the signal to noise ratio is less favorable. It is obvious that in terms of efficiency, sCVD diamonds surpass the performance of pCVD but the commercially available surfaces remain small, which would imply combining several diamonds in the form of a mosaic to make a larger detector.

The purpose of the hodoscope is to detect each incident ion while ensuring 495 intrinsic time resolution ≤ 100 ps. The best results were obtained with the 496 sCVD-DOI reference pair. The TOF resolutions obtained with this pair of 497 detectors are matching the objectives of the project, both with single protons 498 of 68 MeV and with carbon ions of 95 MeV/u. Indeed, the proton & PG 499 TOF resolution obtained during a past ARRONAX experiment [11] showed 500 the capability of our detectors to discriminate PGs with a TOF resolution of 501 101 ps (σ), making techniques such as ultra-fast PGT very promising. Such 502 results could not be obtained with pCVD detectors which exhibit a too low 503 SNR to be able to measure an equivalent timing resolution with 68 MeV pro-504 tons. Moreover, the threshold-based study of their detection efficiency and 505 time resolution demonstrated that combining a detection efficiency > 90%506 and a time resolution at the 100 ps level was not achievable. A noteworthy 507 improvement of their time resolution could only be obtained for threshold 508 values that rejected most of the single-proton signals, thus dramatically de-509 teriorating their detection efficiency. 510

It should be also considered that the energy deposition of a 68 MeV proton is the highest we can get with a single proton in particle therapy. Indeed, the protons energy range varies from 70 MeV to 250 MeV. The deposited energy, and therefore the generated signal, is inversely proportional to the proton's initial energy. The combination of these considerations makes difficult the use of pCVD detectors for time tagging of single protons in the energy range of proton therapy. We will therefore use sCVD detectors, with the limitation on the commercially available area for this application.

However, the results obtained with carbon ions at GANIL are promising. 519 The 13 ps (σ) TOF resolution obtained between the sCVD Element6 detector 520 and the Augsburg DOI one is the best time performance we measured, in 521 all our experiments. This result is mainly explained by the large energy 522 deposition generated by each ion in the diamond and by the quality of the 523 two diamond samples. This energy deposit is so that a 66 ps (σ) resolution 524 between two pCVD detectors was obtained whereas they were metallized 525 with electrodes of 7 and 16 mm in diameter, respectively. Assuming that 526 this value is the quadratic sum of their respective timing resolutions, we can 527 estimate that their individual timing resolution is equivalent to or better 528 than 66 ps. Besides, in the case of carbon ion therapy, the energy of the 520 ions ranges from 95 MeV/u to 400 MeV/u. SRIM simulations show that in a 530 500 µm pCVD diamond with a charge collection efficiency of 30% (measured 531 on an alpha test bench at laboratory) generates a collected charge ranging 532 from 156 fC to 61 fC, respectively (see [36]). As a comparison, a 5.49 MeV 533 α particle (equivalent to 67 fC) generates a sufficient signal to measure an 534 intrinsic resolution of less than 100 ps [21, 22]. We can thus reasonably 535 assume that to obtain an intrinsic temporal resolution of 100 ps (σ) with a 536 large size (pCVD) detector remains a realistic goal for carbon ion therapy. 537

It should also be noted that in the configuration of our tests, pCVD sen-538 sors were not optimized for timing measurements. An improvement of their 539 timing performance could be obtained by combining different approaches. 540 First, their thickness could be reduced down to their charge collection dis-541 tance so that the applied electric field can be higher. By doing so, a seg-542 mentation of the active surface would be necessary to over-compensate the 543 increase of the capacitance. Using two layers of thin pCVD sensors would 544 allow to improve the timing performance of the device by a factor $\sqrt{2}$ and 545 they could be inclined with respect to the beam axis to increase their effective 546 thickness. 547

Finally, concerning the particle counting performance, the measurements 548 carried out with 68 MeV protons at a beam intensity of ~ 6 pA can allow 549 us to conclude that a beam monitor equipped with sCVD diamond sensors 550 makes it possible to provide both fast timing and counting of protons inside 551 a bunch. In terms of hadon herapy beam monitoring, this makes it pos-552 sible to count at the start of treatment at reduced beam intensity and, if 553 necessary, identify bunches where the proton multiplicity is greater than 1. 554 On the contrary, pCVD detectors are not able to achieve particle counting 555 at low proton rate. This result on the comparative performance of sCVD 556 and pCVD diamonds should however be qualified. Indeed, for higher beam 557 intensity, sCVD diamond sensor thickness is certainly to be optimized to 558 prevent long time drift which may result in a pile-up phenomenon at highest 559 RF frequencies (up to 106 MHz). pCVD may present an advantage relative 560 to sCVD. Since charge trapping occurs while charge carriers are drifting to 561 the electrodes, it results in a shorter signal as observed in Figure 9 at ~ 2 nA 562



Figure 9: Compared time-domain responses of the Element6 sCVD and the DDK pCVD detectors, irradiated with the ARRONAX proton beam at $I_{beam} \sim 2 \,\mathrm{nA}$ (with an accelerator radio-frequency of 30.45 MHz). The induced currents produced by the detectors are converted into voltage signals through a 50 Ω resistor.

(~400 protons/bunch at 30.45 MHz). Such a beam current is close to clinical
conditions. Therefore, the two types of diamond could be used depending on
the targeted intensity range.

566 5. Conclusions

The present results are encouraging the development of a beam-tagging hodoscope with TOF capabilities. For all the tests presented in this work, sCVD diamond detectors demonstrated characteristics that are in good agreement with the requirements of the hodoscope project. The detection effi-

ciency measurements highlighted that pCVD detectors can detect single ions 571 with a good efficiency but can not reach a timing resolution at the order of 572 100 ps (σ) when detecting single protons. At low intensity, their poor energy 573 resolution prevent them from counting the number of protons contained in 574 a bunch but their mean charge response remains linear with the deposited 575 energy. At higher intensity, the shorter pulses generated by pCVD detectors 576 can represent an advantage over sCVD for beam monitoring at 100 MHz 577 rates. Using carbon ions, both sCVD and pCVD demonstrated excellent 578 timing results. 579

Consequently, two solutions can be foreseen for the beam tagging ho-580 doscope design. The first one may consist in using either four $4.5 \times 4.5 \times$ 581 0.5 mm^3 commercially available sCVD diamonds arranged in mosaic or, later 582 on, large area sCVD diamonds. The second solution may consist in using 20 583 $\times 20 \times 0.3 \text{ mm}^3$ pCVD mainly dedicated for carbon ion therapy applications. 584 In both cases, the hodoscope will be made out of double-sided strip sensors. 585 It will provide the ion transverse position with a precision $\leq 1 \,\mathrm{mm}^2$ (X and 586 Y strips width). The influence of the segmentation of the metallic contacts 587 on the timing performance of the device will have to be evaluated while it 588 will not be possible to use the side-to-side signal summation method. The 589 next step of the hodoscope development is the assembling of the two selected 590 diamonds types with front-end electronics currently developed at LPSC for 591 TOF measurements of prompt-gamma in view of range verification in particle 592 therapy. 593

594 Acknowledgements

The authors would like to acknowledge the ESRF-ID21 beamline for pro-595 vision of synchrotron radiation with experiments MI-1243 (2016) and MI-596 1285 (2017), and support from the ESRF BCU group for integrating the 597 triggered readout of the LeCroy DSO into the ID21 SPEC data acquisition 598 system. This work was supported by Plan Cancer (CLaRyS-UFT project), 599 the LabEx PRIMES (ANR-11-LABX-0063), FranceHadron (ANR-11-INBS-600 0007) and ANR MONODIAM-HE (ANR-089520). The cyclotron Arronax 601 is supported by CNRS, Inserm, INCa, the Nantes University, the Regional 602 Council of Pays de la Loire, local authorities, the French government and the 603 European Union. This work has been, in part, supported by a grant from 604 the French National Agency for Research called "Investissements d'Avenir", 605 Equipex Arronax-Plus noANR-11-EQPX-0004, Labex IRON noANR-11-LABX-606 18-01and ISITE NExT no ANR-16-IDEX-007. It was performed in the frame 607 of ENSAR2/MediNet network (Horizon2020-654002). The authors are grate-608 ful to Matthias Schreck from Augsburg University and Martin Fischer from 609 Audiatec Augsburg for providing the LPSC laboratory with DOI samples. 610 Dominique Breton and Jihanne Maalmi from IJC-Lab Orsay and Eric De-611 lagnes from CEA Saclay are thanked for their implication in dedicated soft-612 ware development and technical support of the WaveCatcher data acquisition 613 system. SC, MLGM, AB, JC, DD, LGM, AG, AL, SM, OR, FER, and MY 614 are members of the RD42 collaboration at CERN. The authors would like 615 to thank the reviewer for his/her useful discussion about the enhancement 616 of the timing performance of pCVD sensors that improved the quality of the 617 discussion. 618

619 References

- [1] W. D. Newhauser, R. Zhang, The physics of proton therapy,
 Physics in Medicine and Biology 60 (8) (2015) R155–R209.
 doi:10.1088/0031-9155/60/8/R155.
- URL http://stacks.iop.org/0031-9155/60/i=8/a=R155?key=
 crossref.e17ea27b3e09d2ae08a7471562523fb1
- [2] D. Schardt, T. Elsässer, D. Schulz-Ertner, Heavy-ion tumor therapy:
 Physical and radiobiological benefits, Rev. Mod. Phys. 82 (2010) 383–
 425. doi:10.1103/RevModPhys.82.383.
- URL https://link.aps.org/doi/10.1103/RevModPhys.82.383
- [3] H. Paganetti, Range uncertainties in proton therapy and the role of
 Monte Carlo simulations., Physics in Medicine and Biology 57 (11)
 (2012) R99—-117. doi:10.1088/0031-9155/57/11/R99.
- ⁶³² URL http://dx.doi.org/10.1088/0031-9155/57/11/R99
- [4] A. C. Knopf, A. Lomax, In vivo proton range verification: A re view, Physics in Medicine and Biology 58 (15) (2013) 131–160.
 doi:10.1088/0031-9155/58/15/R131.
- ⁶³⁶ URL http://stacks.iop.org/0031-9155/58/i=15/a=R131?key=
 ⁶³⁷ crossref.a4dce585277cdd2c3b0331cb1d3e7322
- [5] A. C. Kraan, Range verification methods in particle therapy: Underlying physics and Monte Carlo modelling, Frontiers in Oncology 5 (JUN)
 (2015) 150. doi:10.3389/fonc.2015.00150.

- [6] J. Krimmer, D. Dauvergne, J. M. Létang, Testa, Prompt-gamma monitoring in hadrontherapy: A review, Nuclear Instruments and Methods
 in Physics Research, Section A: Accelerators, Spectrometers, Detectors
 and Associated Equipment 878 (2018) 58–73. doi:10.1016/j.nima.
 2017.07.063.
- [7] T. Werner, J. Berthold, F. Hueso-González, T. Koegler, J. Petzoldt,
 K. Roemer, C. Richter, A. Rinscheid, A. Straessner, W. Enghardt,
 G. Pausch, Processing of prompt gamma-ray timing data for proton
 range measurements at a clinical beam delivery, Physics in Medicine
 and Biology 64 (10) (2019) 105023. doi:10.1088/1361-6560/ab176d.
 URL https://iopscience.iop.org/article/10.1088/1361-6560/
 ab176d
- [8] O. Allegrini, J.-P. Cachemiche, C. Caplan, B. Barlus, X. Chen, S. Curtoni, D. Dauvergne, R. Della Negra, M.-L. Gallin-Martel, J. Hérault,
 J.-M. Létang, C. Morel, E. Testa, Y. Zoccarato, Characterization of a beam tagging hodoscope for hadrontherapy monitoring, Journal of InstrumentationAccepted manuscript (2020).
- [9] C. Golnik, F. Hueso-González, A. Müller, P. Dendooven, W. Enghardt,
 F. Fiedler, T. Kormoll, K. Roemer, J. Petzoldt, A. Wagner, G. Pausch,
 Range assessment in particle therapy based on prompt γ-ray timing
 measurements, Physics in Medicine and Biology 59 (18) (2014) 5399–
 5422. doi:10.1088/0031-9155/59/18/5399.
- 663 URL http://stacks.iop.org/0031-9155/59/i=18/a=5399?key= 664 crossref.5437fcd3059992135ec2113679c7dad6

- [10] F. Hueso-González, W. Enghardt, F. Fiedler, C. Golnik, G. Janssens,
 J. Petzoldt, D. Prieels, M. Priegnitz, K. E. Römer, J. Smeets, F. Vander
 Stappen, A. Wagner, G. Pausch, First test of the prompt gamma ray
 timing method with heterogeneous targets at a clinical proton therapy
 facility, Physics in Medicine and Biology 60 (16) (2015) 6247–6272.
 doi:10.1088/0031-9155/60/16/6247.
- URL http://stacks.iop.org/0031-9155/60/i=16/a=6247?key=
 crossref.3382b95c39af8f8ab69e65cd74102dff
- [11] S. Marcatili, J. Collot, S. Curtoni, D. Dauvergne, J.-Y. Hostachy,
 C. Koumeir, J. M. Létang, J. Livingstone, V. Métivier, L. Gallin-Martel,
 M. L. Gallin-Martel, J. F. Muraz, N. Servagent, É. Testa, M. Yamouni,
 Ultra-fast prompt gamma detection in single proton counting regime for
 range monitoring in particle therapy, Physics in Medicine & Biology
 65 (24) (2020) 245033. doi:10.1088/1361-6560/ab7a6c.
- ⁶⁷⁹ URL https://doi.org/10.1088/1361-6560/ab7a6c
- [12] D. Dauvergne, O. Allegrini, C. Caplan, X. Chen, S. Curtoni,
 A. Etxebeste, M.-L. Gallin-Martel, M. Jacquet, J. M. Létang, J. Livingstone, S. Marcatili, C. Morel, E. Testa, Y. Zoccarato, On the Role
 of Single Particle Irradiation and Fast Timing for Efficient OnlineControl in Particle Therapy, Frontiers in Physics 8 (2020) 434. doi:
 10.3389/fphy.2020.567215.
- ⁶⁸⁶ URL https://www.frontiersin.org/article/10.3389/fphy.2020.
 ⁶⁸⁷ 567215
- ⁶⁸⁸ [13] A. Vignati, V. Monaco, A. Attili, N. Cartiglia, M. Donetti, M. F. Maz-

689	inani, F. Fausti, M. Ferrero, S. Giordanengo, O. H. Ali, M. Mandur-
690	rino, L. Manganaro, M. Ferrero, G. Mazza, R. Sacchi, V. Sola, A. Sta-
691	iano, R. Cirio, Innovative thin silicon detectors for monitoring of thera-
692	peutic proton beams: preliminary beam tests, Journal of Instrumenta-
693	tion 12 (12) (2017) C12056–C12056. doi:10.1088/1748-0221/12/12/
694	c12056.
695	URL https://doi.org/10.1088%2F1748-0221%2F12%2F12%2Fc12056
696 [14]	L. Federici, G. Aglieri Rinella, D. Alvarez Feito, R. Arcidiacono,
697	C. Biino, S. Bonacini, A. Ceccucci, S. Chiozzi, E. Cortina Gil,
698	A. Cotta Ramusino, J. Degrange, M. Fiorini, E. Gamberini, A. Gianoli,
699	J. Kaplon, A. Kleimenova, A. Kluge, A. Mapelli, F. Marchetto,
700	E. Migliore, E. Minucci, M. Morel, J. Noël, M. Noy, L. Perktold,
701	M. Perrin-Terrin, P. Petagna, F. Petrucci, K. Poltorak, G. Romagnoli,
702	G. Ruggiero, B. Velghe, H. Wahl, The Gigatracker, the silicon beam
703	tracker for the NA62 experiment at CERN, Nuclear Instruments
704	and Methods in Physics Research Section A: Accelerators, Spec-
705	trometers, Detectors and Associated Equipment 958 (2020) 162127,
706	proceedings of the Vienna Conference on Instrumentation 2019.
707	doi:https://doi.org/10.1016/j.nima.2019.04.081.
708	URL http://www.sciencedirect.com/science/article/pii/

- 709 S0168900219305637
- [15] E. Bossini, N. Minafra, Diamond Detectors for Timing Measurements
 in High Energy Physics, Frontiers in Physics 8 (2020) 248. doi:
 10.3389/fphy.2020.00248.

URL https://www.frontiersin.org/article/10.3389/fphy.2020. 00248

- [16] M. Pomorski, E. Berdermann, A. Caragheorgheopol, M. Ciobanu,
 M. Kiš, A. Martemiyanov, C. Nebel, P. Moritz, Development of singlecrystal CVD-diamond detectors for spectroscopy and timing, Physica
 Status Solidi (A) Applications and Materials Science 203 (12) (2006)
 3152–3160. doi:10.1002/pssa.200671127.
- ⁷²⁰ [17] Element6, https://e6cvd.com//application/quantum-radiation.html.
- ⁷²¹ [18] II-VI Inc., https://ii-vi.com/product/cvd-diamond-substrates/.
- ⁷²² [19] US Applied Diamond Inc., http://usapplieddiamond.com/products/.
- [20] Augsburg diamond technology gmbh (audiatec),
 https://www.audiatec.de/.
- [21] M. L. Gallin-Martel, A. Bes, A. Boukhémiri, G. Bosson, J. Col-725 lot, D. Dauvergne, M. Fontana, L. Gallin-Martel, A. Gorecki, J. Y. 726 Hostachy, J. Krimmer, A. Lacoste, S. Marcatili, J. Morse, J. F. Mu-727 raz, F. E. Rarbi, O. Rossetto, M. Salomé, E. Testa, M. Yamouni, 728 Large area polycrystalline diamond detectors for online hadron ther-729 apy beam tagging applications, in: 2016 IEEE Nuclear Science Sympo-730 sium, Medical Imaging Conference and Room-Temperature Semicon-731 ductor Detector Workshop (NSS/MIC/RTSD), 2016, pp. 1-5. doi: 732 10.1109/NSSMIC.2016.8069398. 733
- ⁷³⁴ URL https://doi.org/10.1109/NSSMIC.2016.8069398

735	[22]	M. L. Gallin-Martel, L. Abbassi, A. Bes, G. Bosson, J. Collot, T. Crozes,
736		S. Curtoni, D. Dauvergne, W. De Nolf, M. Fontana, L. Gallin-Martel,
737		J. Y. Hostachy, J. Krimmer, A. Lacoste, S. Marcatili, J. Morse, J. F.
738		Motte, J. F. Muraz, F. E. Rarbi, O. Rossetto, M. Salomé, Testa,
739		R. Vuiart, M. Yamouni, A large area diamond-based beam tagging
740		hodoscope for ion therapy monitoring, in: EPJ Web of Conferences,
741		Vol. 170, EDP Sciences, 2018, p. 09005. doi:10.1051/epjconf/
742		201817009005.

[23] A. Lacoste, T. Lagarde, S. B. chu, Y. Arnal, J. Pelletier, Multidipolar plasmas for uniform processing: physics, design and performance, Plasma Sources Science and Technology 11 (4) (2002) 407–412.
doi:10.1088/0963-0252/11/4/307.

⁷⁴⁷ URL https://doi.org/10.1088%2F0963-0252%2F11%2F4%2F307

- ⁷⁴⁸ [24] CIVIDEC Instrumentation, https://cividec.at/.
- [25] D. Breton, E. Delagnes, J. Maalmi, P. Rusquart, The WaveCatcher
 family of SCA-based 12-bit 3.2-GS/s fast digitizers, in: 2014 19th IEEENPSS Real Time Conference, RT 2014 Conference Records, Institute
 of Electrical and Electronics Engineers Inc., 2015. doi:10.1109/RTC.
 2014.7097545.
- F. Poirier, S. Girault, S. Auduc, C. Huet, E. Mace, J. L. Delvaux,
 F. Haddad, The C70 ARRONAX and beam lines status, in: IPAC 2011 2nd International Particle Accelerator Conference, 2011, pp. 2661–2663.
- 757 [27] F. Poirier, S. Girault, F. B. Harel, J. B. Etienne, X. Goiziou,

- F. Gomez, A. Herbert, L. Lamouric, D. Poyac, H. Trichet, C. Huet,
 E. Mace, Studies and Upgrades on the C70 Cyclotron Arronax, in:
 Proceedings of Cyclotrons 2016, 2016, pp. 235–237. doi:10.18429/
 JACoW-Cyclotrons2016-TUD02.
- URL http://jacow.org/cyclotrons2016/papers/tud02.pdf
- [28] P. Moritz, E. Berdermann, K. Blasche, H. Stelzer, B. Voss, Broadband
 electronics for CVD-diamond detectors, Diamond and Related Materials
 10 (9-10) (2001) 1765–1769. doi:10.1016/S0925-9635(01)00434-4.
- [29] M. Berretti, E. Bossini, N. Minafra, Timing performance of diamond
 detectors with Charge Sensitive Amplifier readout, Tech. Rep. September, CERN (2015).
- 769 URL http://cds.cern.ch/record/2055747?ln=fr
- [30] M. Cotte, E. Pouyet, M. Salomé, C. Rivard, W. De Nolf, H. Castillo-770 Michel, T. Fabris, L. Monico, K. Janssens, T. Wang, P. Sciau, L. Verger, 771 L. Cormier, O. Dargaud, E. Brun, D. Bugnazet, B. Fayard, B. Hesse, 772 A. E. Pradas del Real, G. Veronesi, J. Langlois, N. Balcar, Y. Van-773 denberghe, V. A. Solé, J. Kieffer, R. Barrett, C. Cohen, C. Cornu, 774 R. Baker, E. Gagliardini, E. Papillon, J. Susini, The ID21 x-ray and 775 infrared microscopy beamline at the ESRF: status and recent applica-776 tions to artistic materials, J. Anal. At. Spectrom. 32 (2017) 477–493. 777 doi:10.1039/C6JA00356G. 778
- ⁷⁷⁹ URL http://dx.doi.org/10.1039/C6JA00356G
- [31] M.-L. Gallin-Martel, S. Curtoni, S. Marcatili, L. Abbassi, A. Bes,
 G. Bosson, J. Collot, T. Crozes, D. Dauvergne, W. De Nolf,

M. Fontana, L. Gallin-Martel, A. Ghimouz, J.-Y. Hostachy, A. Lacoste, J. Morse, J.-F. Motte, J.-F. Muraz, F. Rarbi, O. Rossetto, M. Salomé, E. Testa, M. Yamouni, X-ray Beam Induced
Current analysis of CVD diamond detectors in the perspective of a beam tagging hodoscope development for hadrontherapy on-line monitoring, Diamond and Related Materials (2020)
108236doi:https://doi.org/10.1016/j.diamond.2020.108236.

- ⁷⁸⁹ URL http://www.sciencedirect.com/science/article/pii/
 ⁷⁹⁰ S0925963520307913
- [32] M. Berger, J. Hubbell, S. Seltzer, J. Chang, J. Coursey, R. Sukumar,
 D. Zucker, K. Olsen, XCOM: Photon Cross Section Database (2010).
 doi:https://dx.doi.org/10.18434/T48G6X.
- ⁷⁹⁴ URL https://www.nist.gov/pml/xcom-photon-cross-sections-database
- [33] H. Frais-Kölbl, E. Griesmayer, H. Kagan, H. Pernegger, A fast low-noise
 charged-particle CVD diamond detector, IEEE Transactions on Nuclear
 Science 51 (6 III) (2004) 3833–3837. doi:10.1109/TNS.2004.839366.
- [34] M. Ciobanu, E. Berdermann, N. Herrmann, K. D. Hildenbrand,
 M. Kiš, W. Koenig, J. Pietraszko, M. Pomorski, M. Rebisz-Pomorska,
 A. Schüttauf, In-beam diamond start detectors, IEEE Transactions on
 Nuclear Science 58 (4 PART 2) (2011) 2073–2083. doi:10.1109/TNS.
 2011.2160282.
- [35] J. F. Ziegler, M. D. Ziegler, J. P. Biersack, SRIM The stopping and
 range of ions in matter (2010), Nuclear Instruments and Methods in
 Physics Research, Section B: Beam Interactions with Materials and

- Atoms 268 (11-12) (2010) 1818-1823. doi:10.1016/j.nimb.2010.02. 807 091.
- [36] S. Curtoni, Development of a diamond beam-tagging hodoscope demonstrator for online ion range verification in hadrontherapy, Ph.D. thesis,
 Université Grenoble-Alpes (2020).