Donation after circulatory death program in Italy

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Abstract: Donation after cardiac death (DCD) donors may help increase the donor pool for lung transplantation. Here, we briefly describe the Italian organ transplantation system and present the Italian DCD program. Our country adopts a mixed “opting-in” and “opting-out” system. Death declaration is confirmed by neurological or cardiocirculatory criteria; in case of cardiac death, the Italian law requires 20 minutes of documented asystole. Organs are primarily allocated to urgent patients, otherwise they are offered to a centre within the region. Lung transplantation centres in Lombardy, instead, use Lung Allocation Score (LAS). In Italy organ donation activity and transplantation has been growing, but the gap remains between patients on the waiting list and the number of transplantations. DCDs may alleviate donor shortage, but the path towards a DCD Italian program was complicated, and physicians had to face the challenge of organ preservation with a prolonged no-touch period. The first DCD program (Alba protocol) started in 2007 and proved DCD after unexpected cardiac arrest [uncontrolled DCD (uDCD)] possible for kidney transplantation, using post-mortem normothermic regional perfusion (NRP) before recovery. In 2015 the first DCD liver transplantation in Italy was performed at Niguarda Hospital in Milan using innovative strategies based on NRP and ex-situ organ perfusion. The Careggi Teaching Hospital in Florence started a DCD protocol for kidney and liver transplantation. The first lung transplantation from an uDCD donor in Italy was performed at the Policlinico transplant centre in Milan in 2014: our protocol consists of a normothermic open-lung preservation, namely without chest drainages for topical cooling, avoiding lung hypoxia through recruitment manoeuvres, continuous positive airway pressure (CPAP), and protective ventilation. Lungs are assessed using ex-vivo lung perfusion (EVLP). Eventually, we also began including controlled DCDs (cDCDs). A dedicated protocol for thoracic and abdominal organs retrieval was established in 2017, combining NRP with our open lung approach, with good results. Over the last 5 years, transplantation with grafts from DCDs has been increasing: in 2019, they represented 4.5% of such procedures and 8.5% of lung transplantations. Our results showed the feasibility of combined procurement in different settings with no detrimental effects on abdominal organs despite extended ischemia times.

Keywords: Donation after cardiac death (DCD); transplantation; lung transplantation; Italy; organ allocation

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Lung transplantation is a consolidated clinical reality and has become a worldwide established procedure for selected patients suffering from end stage pulmonary failure. In time, the number of such procedures has been increasing, showing substantial improvement in both patients’ survival and quality of life. However, these outcomes are still affected by both early and late complications (1). Nevertheless, significant differences among countries exist, mostly with regard to organ donation and donor management. Culture, ethics, and law seriously affect these aspects; surgical techniques and recipient postoperative care, instead, appear more homogeneous.

The Italian donation system

Consent to donation and death declaration

In Italy, a presumed consent law for organ donation was approved in 1999, stating that citizens were to express their willingness or refusal to donate organs after death. Their choice is then recorded in the national organ donor registry. By law, the absence of such declaration is considered consent (2). However, this law has never been enforced, and a mixed “opting-in” and “opting-out” system has been applied to date. Organ and tissue recovery are only possible after death declaration and after confirming the willingness to donate in the national registry, or, in absence of an expressed preference, after obtaining the non-opposition by the donor’s next of kin. According to Italian law, death is defined as the irreversible loss of brain function (3); the determination criteria, both neurological and cardio-circulatory, were established by a decree (4) in order to guarantee the observance of the dead donor rule: organ recovery must not cause the death of the donors, who must be dead before the beginning of organs procurement. For adult patients showing encephalic lesions who are in a state of unconsciousness, have neither brainstem nor respiratory reflexes, and whose electroencephalogram (EEG) shows electro-cerebral silence, brain death is confirmed after 6 hours. During this observation period, the patient is tested at least twice to confirm the loss of brain and brain-stem functions (i.e., absence of reflexes, hypercapnia and acidosis, flatline EEG). Cardiac death ascertainment is based on the cessation of cardio-circulatory functions: according to Italian law, 20 minutes of flat electrocardiogram (ECG) must be recorded in order to declare the patient’s death, as it testifies to the total loss of cerebral functions. In both cases, death is ultimately defined by the cessation of brain activity: with neurological criteria the loss of brain function is directly demonstrated, while in case of cardiac death it is guaranteed by the prolonged lack of cerebral blood flow.

Lung allocation system

In Italy, there are ten lung transplant centres, including one managing paediatrics only, coordinated by a well-established national and regional transplantation network (Figure 1). When a potential donor is referred by the intensive care unit (ICU) staff to the regional coordinating centre [Centro Regionale Trapianti (CRT)], lungs are primarily allocated to urgent patients included in the Italian Urgent Lung Transplant (IULT) program by the National Transplantation Centre [Centro Nazionale Trapianti (CNT)], the national coordinating centre. The IULT program was instituted in 2010: urgent patients are put on a single national waiting list. The priority of each patient lasts 1 week and can be renewed twice. The IULT program inclusion and exclusion criteria are shown in Table 1 (6). If
there are no urgent recipients, organs are allocated within the region. As opposed to the other Italian regions, where there is only one lung transplant centre, in Lombardy, the most densely populated in northern Italy (422,76 inhabitants per square kilometres), there are three lung transplantation centres, including our own. In our region, since March 15\textsuperscript{th}, 2016, lung allocation is no longer “centre oriented” but “patient oriented”, as Lung Allocation Score (LAS) has been applied. The positive experiences in the United States first and then in Germany led the Lombardy Regional Reference Centre to merge the three centres waiting lists and implement LAS for the allocation of lungs within the region (7). Finally, it should be pointed out that Italy’s elongated and narrow morphology (Figure 1), its great number of mountains, covering 40\% of the country’s territory, and the presence of two large islands (Sicily and Sardinia) make rapid and timely transportation of the retrieval team and the grafts quite complicated. These peculiar geographical features might be the cause of very long transfer times for relatively short distances.

### Italian transplantation experience

During the last 20 years, the number of potential donors referred by ICUs and consequently the organ donation activity in general, has been progressively growing (Figure 2). However, as the amount of lung transplantation procedures performed in Italy every year has almost tripled, increasing from 60 lung transplantation in 2000 to 153 in 2019, the number of patients on the waiting list has also slowly yet significantly been rising. The average time period spent on the waiting list for lung transplantation, according to the 2019 CNT report, was 2.5 years (5). At the time, the most severe limitation was, and still is, the shortage of suitable donors. The dramatic consequence of the gap between supply and demand of donor graft was high mortality for patients on the waiting list. The graph shows the gap between the patients on the waiting list and the number of transplants per year in the last 5 years (Figure 3).

Several strategies have been implemented to overcome this shortage, such as the use of extended criteria donors (ECDs) and the donation after circulatory determination of death.

### Table 1 Italian Urgent Lung Transplant (IULT) program inclusion criteria

<table>
<thead>
<tr>
<th>IULT program inclusion criteria</th>
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<tr>
<td>Patients on the waiting list dependent on mechanical ventilation and/or extracorporeal membrane</td>
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<tr>
<td>oxygenator (except for Extracorporeal Carbon Dioxide Removal device)</td>
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<tr>
<td>Patients hospitalized at the transplant centre</td>
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<tr>
<td>Age ≤50 years old</td>
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<tr>
<td>Absence of sepsis, multi-organ failure, haemorrhagic shock, neurologic impairment</td>
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![Figure 2](http://dx.doi.org/10.21037/ccts-20-116)

**Figure 2** Utilised donors per million population (all organs) (5).
Over the years, particularly at the beginning of the lung transplantation era, only “ideal” lung donor grafts were considered for transplantation. However, most of the potential donors in ICUs do not even meet the criteria defining a “standard” lung donor (Table 2). In fact, despite careful management, the prolonged mechanical ventilation in the ICU damages the lungs and the other organs. In Italy, only 40% of referred donors are suitable for multiorgan procurement, and, barely 15–20% of them are considered for lung transplantation, also due to their advanced age. Protocols regarding potential donors’ management have been applied in Italian ICUs in order to maintain the donor stability and optimize the organs’ condition. At the same time, transplantation centres started pushing the limits by accepting organs from donors with increasingly extended criteria. For this reason, research started focusing on finding a way to thoroughly and safely evaluate these “extended criteria” grafts.

### Ex-vivo lung perfusion (EVLP)

The advent of EVLP provided a fundamental tool for lung transplantation centres, allowing for detailed assessment and reconditioning of lungs. In Italy, machine perfusion protocols vary among centres (8,9). In our centre in Milan, we perform a 4-hour-long normothermic EVLP with an open atrium technique, using a perfusion solution made of both Steen solution and red blood cells (reaching a 5–10% haematocrit) and maintaining the target flow at 40% of the estimated donor’s cardiac output, as previously described (Figure 4) (10). Our clinical experience with machine perfusion began in 2011, when our team performed the first lung transplantation after EVLP in Italy, and we have been using it ever since, with good results (11). In 2016, we started using portable organ care system (OCS) for selected cases as well. EVLP proved a valuable resource, as it helped increase the number of lung transplantations performed in our centre, it allowed us to assess and transplant lungs from donors on extra-corporeal membrane oxygenation (ECMO) support and to act as a “lung repair centre” (12). Furthermore, EVLP opened up the possibility for our centre to implement a DCD lung transplantation program.

### DCD donors

DCD describes the retrieval of organs following death, according to circulatory instead of neurological criteria. The definition of death and the dead donor rule remain the same and must be respected. The first lung transplantations

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**Table 2** Standard donor selection criteria for lung transplantation

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<tr>
<td>ABO compatibility</td>
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<tr>
<td>PaO₂ &gt;300 (FiO₂ =1.0; PEEP =5 cmH₂O)</td>
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<tr>
<td>Age &lt;55 years old</td>
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<tr>
<td>Absence of infiltrates at chest X-ray</td>
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<tr>
<td>Absence of secretions/aspiration signs at bronchoscopy</td>
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<tr>
<td>No history of cardiothoracic surgery</td>
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<tr>
<td>Adequate size match</td>
</tr>
<tr>
<td>Smoking history &lt;20 pack/years</td>
</tr>
<tr>
<td>No trauma of the chest</td>
</tr>
</tbody>
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PaO₂, arterial oxygen pressure; FiO₂, fraction of inspired oxygen; PEEP, positive end-expiratory pressure.
Around the world, experiences with DCD are extremely diversified. Notably, some countries do not allow organ DCD (i.e., Germany). The differences among countries mainly depend on local factors regarding culture, logistics, law, ethics and religion. To this, we must add the graft factors, as each organ has its peculiar requirements. All these variables lead to different solutions in DCD approaches as well as preservation and procurement techniques. The “hands off” time is 5 minutes for almost all centres around the world except for Zurich (10 minutes), some Australian centres (2 minutes) and Italy (notably, 20 minutes). Also, in some countries (such as the United Kingdom, Ireland and the Netherlands), ante mortem drug administration aiming at organ preservation is not allowed, while in other centres (for example in Belgium, Spain, Austria and France) it is legal to do so once the next of kin has been informed.

**DCD donors in Italy**

In Italy, heparin can only be given to the donor during the agonal phase, and cannulation is not allowed before the diagnosis of death, whilst femoral vessels wires can be positioned to simplify the upcoming cannulation. Essentially, in Italy it is legally allowed to retrieve organs from DCD donors; from a bioethical perspective, it should be emphasized that the current regulations seek to avoid disregarding the person’s will to donate their organs after death. On the other hand, the path to include DCD donors in clinical practice was complex and far from straightforward. In fact, it should be pointed out that transplant centres in Italy had to face the challenge of an extremely long warm ischemic “no-touch” period of 20 minutes, making organ preservation particularly demanding. In Italy, an uDCD program started in 2007, while the use of controlled donors was only introduced later in 2015 as a direct consequence of the previously mentioned different ethical issues between the two settings. The first uDCD program for kidney transplantation, the “Alba program”, was started in Pavia: according to this protocol, immediately after determination of death with circulatory criteria, ECMO support and then selective abdominal normothermic venous-arterial circulation is established (17). At first, this project had to face some major challenges, including involvement and cooperation with emergency services and the need for on-call dedicated DCD task force personnel. However, despite the long no-touch period required by the Italian legislation, the Alba program had the merit to show that uDCD is feasible even

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**Classification**

In 1995 in Maastricht, the First International Workshop on DCD drafted four categories of patients in order to outline the approach to this type of donors (16); the classification has undergone some changes since its original inception, but the fundamental concepts have remained the same. Specifically, it is possible to identify two settings, uncontrolled and controlled: the first two categories include donors qualified as uncontrolled DCD (uDCD), because cardiac arrest occurs unexpectedly. Additionally, the clinical history of the donor is unknown. In this setting, the greatest difficulties concern the logistical and organisational aspects, as well as the need to thoroughly evaluate the organs. Conversely, we refer to those in the third category as controlled DCD (cDCD) donors, since the procurement is substantially planned after an anticipated cardiac arrest, and donors are largely evaluated in time. For category III donors, the most relevant problem is of ethical nature, and it is related to the suspensions of treatment. Category IV includes brain-dead donors who suffer an unexpected cardiac arrest after death declaration and before organ retrieval has been planned. The modified Maastricht classification of DCD also includes a fifth category, comprising organ donation after euthanasia, which is illegal in Italy.

**Differences around the world**

Around the world, experiences with DCD are extremely
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in our country. The second DCD program was started in Florence at the Careggi Teaching Hospital and managed to transplant both kidneys and liver (18). In 2015, the first successful DCD liver transplantation in Italy was performed at the Niguarda transplant centre in Milan. A DCD program has been active ever since. Normothermic regional perfusion (NRP) was used to assess and preserve abdominal organs, counteracting the prolonged warm ischemia time (19).

The Milan lung transplant centre’s experience

The Policlinico transplant centre in Milan has been a pioneer in lung transplantation with grafts from DCD donors. Our Lung-DCD program started in 2014 and was preceded by a long pre-clinical and an educational phase; at first, we focused on uDCD donors. Our protocol, which was designed for lung procurement only, was based on the observation that lung tissue can dissociate ischaemia from hypoxia, hence the possibility to preserve lungs for an extended period of time by using continuous positive airway pressure (CPAP). Recruitment manoeuvres were performed following diagnosis of death and CPAP was employed. After obtaining consent by the next of kin, the donor was administered heparin and cardiopulmonary resuscitation (CPR) was resumed for 3 minutes. Lungs were ventilated during procurement, and subsequently evaluated using EVLP (Figure 5) (20). In 2014 we performed the first lung transplantation from an uDCD donor in Italy: the recipient is still alive and in good condition. Our protocol differs from the one described by Steen and used by the Spanish group, as we do not insert chest tubes for topical cooling, leaving the body intact until the procurement in the operating theatre, at virtually no costs. This peculiar strategy allowed us to apply this protocol in both central hospitals with an extracorporeal life-support (ECLS) program, and less-equipped peripheral hospitals. Also, it guarantees an adequate time to face organisational

Figure 5 Timeline for uDCD lung retrieval according to our protocol. uDCD, uncontrolled donation after cardiac death; OOH, out of hospital; ED, Emergency department; CA, cardiac arrest; CPR, cardiopulmonary resuscitation; EKG, electrocardiogram; DOD, declaration of death; NOK, next of kin; ev, endovenous; RM, recruitment manoeuvres; FOB, flexible fiberoptic bronchoscopy; rTPA, recombinant tissue plasminogen activator; EVLP, ex-vivo lung perfusion; WIT, warm ischemia time; CIT, cold ischemia time. cPAP, continuous positive airway pressure; FiO2, fraction of inspired oxygen; RR, respiratory rate; TV, tidal volume; PEEP, positive end-expiratory pressure; I/E, inspiration/expiration.
issues during in-situ preservation and an adequate ex-vivo evaluation. After gaining some experience with the uncontrolled setting, we eventually decided to explore other options by including cDCDs in our project as well. In fact, at that time the concept of withdrawal of life-sustaining treatments (WLSTs) when deemed futile, was beginning to spread and to be considered ethically suitable around ICUs in Italy, prompting us to adapt our project to these different settings. On the other hand, the growing experience on abdominal organs transplantation from DCD donors throughout Italian centres urged us to cooperate in order to establish a dedicated combined procurement protocol, which was drawn up by a multidisciplinary team (21). The required 20 minutes of no-touch posed a challenge for liver retrieval, as it inevitably suffers from a prolonged warm ischemia time: this obstacle was overcome by using NRP. In our protocol we combined the use of NRP for abdominal organ preservation, in order to both minimize warm ischemia damage and assess the graft function, with our normothermic open-lung approach (Figure 6). The NRP stability is tested by closing the inferior vena cava through a tourniquet and clamping the aorta (Figure 7). All organs (lungs, kidneys, liver) are assessed through ex-situ perfusion, as they endure a prolonged functional warm ischemic time. Machine perfusion is considered particularly mandatory for DCD donors in an uncontrolled setting. This Full-DCD project began in October 2017: during the first 2 months, five cDCD donors from four different hospitals were managed according to this protocol. Two of these donors were ECDs, while another one was on ECLS support. In three out of five cases lungs were judged suitable and transplanted at our centre. To this day, all recipients are alive and in good health. In one case the lungs were deemed inadequate for transplantation due to infection, and therefore were not even retrieved; in another case, they showed bronchorrhea at the end of EVLP evaluation and could not be transplanted. Livers were used for transplantation in all five cases, with good outcomes; six out of ten procured kidneys were transplanted. These data showed that there were no detrimental effects on
abdominal organs: performing a double NRP test allowed for a dynamic adjustment of flow settings. These excellent results pushed us to continue our path and reassured the abdominal surgeons. At our centre, from November 1st, 2014 to July 2019, 11 transplantation procedures were carried out with grafts coming from DCD donors; five of those were uDCDs and six were cDCDs. All patients are still alive. Remarkably, our combined retrieval DCD protocol helped increase the number of performed lung transplantation, showing promising results and no reduction in the number of transplanted abdominal organs. Furthermore, procurement from non-heart-beating donors allows us to avoid the cytokine storm and its damage to the organs. However, due to the prolonged ischemic time grafts from DCDs have to endure, we routinely use EVLP in order to assess the organs function. As some studies have already shown that *ex-vivo* evaluation is not mandatory for grafts retrieved from category III DCD donors (22), we are too considering reserving EVLP for uDCDs and for cDCDs when the *in-vivo* assessment demonstrate suboptimal graft function. In general, the transplantation experience with DCD donors has undeniably become relevant in our country. Over the last 5 years, the number of transplantation procedures performed in Italy with grafts from DCD donors has been increasing: in 2019, 115 out of 3,449 transplantations from deceased donors were from DCDs (4.5%). If we only consider lung transplantation, this rate rises to 8.5% and other centres have followed our experience in these years. In conclusion, our protocol offers a very adaptable approach that has been applied in various scenarios: cDCD and uDCD settings, isolated and combined thoracic and abdominal retrieval, hospitals with or without ECLS programs. At the expense of longer ischemic times, we prefer to optimise the lungs’ conditions by normothermic ventilation. By using this strategy, we avoid blindly perfusing the lungs before visual inspection and recruitment, which could result in inhomogeneous cold flushing in case of atelectasis or pleural effusion. It proved safe, as grafts function is assessed both *in-situ* during normothermic ventilation and *ex-situ* during normothermic perfusion, and it demonstrates the feasibility of a combined approach, overcoming the obstacle of prolonged warm ischemic time. The complexity of the program, however, requires a continuous education for the staff involved, especially for the uncontrolled setting. We are currently working on involving a growing number of hospitals in our project. Our experience shows the feasibility of a multi-organ donation from DCD donors, confirming it is possible to abide by the dead donor rule and to overcome the prolonged “no-touch” period required by law.

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Footnote

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